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I, JONNE YABSLEY, TEAM LEADER EXAMINATION SUPPORT AND
SALES hereby certify that annexed is a true copy of the Provisional specification
in connection with Application No. 2002951868 for a patent by PRANA
BIOTECHNOLOGY LIMITED as filed on 04 October 2002.



WITNESS my hand this
Fourteenth day of October 2003

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AUSTRALIA
Patents Act 1990

PROVISIONAL SPECIFICATION

Applicant(s):

PRANA BIOTECHNOLOGY LIMITED

Invention Title:

AZAQUINOLINE DERIVATIVES

The invention is described in the following statement:

AZAQUINOLINE DERIVATIVES

The present invention relates to azaquinoline derivatives, processes for their preparation and their use as pharmaceutical or veterinary agents, in particular for the treatment of neurological conditions, more specifically neurodegenerative conditions such as Alzheimer's disease.

BACKGROUND OF THE INVENTION

All references, including any patents or patent applications, cited in this specification are hereby incorporated by reference. No admission is made that any reference constitutes prior art. The discussion of the references states what their authors assert, and the applicants reserve the right to challenge the accuracy and pertinency of the cited documents. It will be clearly understood that, although a number of prior art publications are referred to herein, this reference does not constitute an admission that any of these documents forms part of the common general knowledge in the art, in Australia or in any other country.

The life span is thought to be biologically fixed for each species, and the length of the human life span is uncertain, but may be up to 120 years. Since life expectancy has risen significantly in this century, the elderly are an increasing segment of our population, and their health care needs will continue to grow for decades. Although normal aging is characterized by modest reductions in the mass and volume of the human brain, which may be due to the atrophy and/or death of brain cells, these changes are far more profound in the brains of patients who succumb to a neurodegenerative condition. Most of these conditions are sporadic (i.e., not due to genetic mutations) and of unknown cause, but hundreds of different mutations in many genes have been shown to cause familial (inherited) variants of several neurodegenerative conditions. Many of the dozen or more genes that harbor these mutations were discovered in the quest to determine the genetic basis of neurodegenerative conditions just in the last ten years. Neurodegenerative conditions evolve gradually after a long period of normal brain function, due to progressive degeneration (i.e., nerve cell dysfunction and death) of specific brain regions. Since symptomatic expression of disease occurs when nerve cell loss exceeds a "threshold" for the continuing function (e.g., memory, movement) performed by the affected brain region, the actual onset of brain degeneration may precede clinical expression by many years.

Intellectual and higher integrative cognitive faculties become progressively impaired and interfere with activities of daily living in neurological

conditions resulting in dementia. The precise prevalence of dementia in the elderly population is unknown, but may be 15% of people over 65 years old with 5% severely and 10% mildly to moderately demented. The prevalence of severe dementia increases from 1% at 65 years to 45% at 85 years. There are many causes of dementia, but

5 Alzheimer's Disease (AD) accounts for 50% of demented patients over 65 years of age.

AD is a primary degenerative disease of the brain. It is characterized by progressive decline of cognitive functions such as memory, thinking, comprehension, calculation, language, learning capacity and judgement. Dementia is diagnosed when these declines are sufficient to impair personal activities of daily living. AD shows an
10 insidious onset with slow deterioration. This disease needs to be clearly differentiated from age-related normal decline of cognitive functions. The normal decline is much less, much more gradual and leads to milder disabilities. The onset of AD is usually after 65 years of age, although earlier onset is not uncommon. As age advances, the incidence increases rapidly (it roughly doubles every 5 years). This has obvious
15 implications for the total number of individuals living with this disorder as life expectancy increases in the population.

The aetiology of dementia of AD is unclear. There is considerable evidence of a heritable predisposition for some forms of AD (reviewed in St George-Hyslop, 2000), and the expression of certain isoforms of ApoE has also been linked to a
20 higher risk of AD (Corder et al, 1993; Czech et al 1994). The toxic accumulation of aluminium has been suggested as a causative agent in AD, although this hypothesis has now been largely superseded. The brains of AD patients display abnormal deposits which include β -amyloid protein ($A\beta$).

$A\beta$ is known to be present in the brains of individuals with certain
25 neurodegenerative diseases, but it is not known whether it is symptomatic of an underlying disease process, or is actually involved in the aetiology of the disease. For example, some authors believe that the $A\beta$ deposits may be indicative of a normal brain defence mechanism, in which the brain attempts to sequester the $A\beta$; such deposits can be present in the brains of normal individuals. There is a mutation of *tau* protein in
30 which neurofibrillary tangles, but no amyloid plaques are present in the brain; this condition is known as tauopathy.

One proposed approach to AD therapy is to inhibit production of $A\beta$ in the brain. Proteolytic cleavage of APP by BACE1 and γ -secretase generates the full-length $A\beta$, which is then released from cells (Nunan and Small, 2000). Therefore
35 inhibitors of either BACE1 or γ -secretase may be of therapeutic value. Alternatively, a number of studies have shown that cholesterol can influence $A\beta$ release (Simons et al., 1998; Hartmann, 2001; Fassbender et al., 2001; Frears et al., 1999; Friedhoff et al.,

2001). Therefore inhibitors of cholesterol biosynthesis, such as statins, may also be of therapeutic value. One advantage of statins is that they have relatively low toxicities, and their mode of action is much better understood than many other compounds currently being investigated as therapeutic agents for AD. However, there is some
5 disagreement in the art as to the value of lowering cholesterol levels, and some workers consider that cholesterol is actually beneficial. For example, Ji et al, (2002) have suggested that the binding of A β to cholesterol might prevent A β toxicity by inhibiting its oligomerization.

A β can bind to lipids (Curtain et al., 2001; Valdez-Gonzalez et al.,
10 2001), including gangliosides (Ariga et al., 2001).

It has been suggested that A β can bind to membrane lipids, and this interaction may be toxic for cells (Hertel et al., 1997). However, few studies have attempted to correlate the degree of lipid binding by A β with its toxicity.

It is also known that acetylcholinesterase (AChE) colocalises with A β in
15 the amyloid deposits which are found in the brains of Alzheimer's disease patients, and that AChE accelerates amyloid formation, both from wild-type A β and from a mutant A β peptide which alone is able to produce few amyloid-like fibres.

In an alternative approach, it has been proposed that by unravelling the proteolytic processing of the amyloid precursor protein (APP), which generates the A β
20 amyloid monomer, a number of possible therapeutic targets may be possible (Shearman et al., 2000; Sinha et al., 1999), and this approach is in an early stage of clinical development. Attempts to promote the clearance of A β from the brain through immunization with A β , while efficacious in a transgenic mouse model for AD (Schenk et al 1999), have been found to have significant adverse effects (Brower, 2002).

25 It has also been suggested that deposition of amyloid-like fibrils may also be important in other neurodegenerative diseases. These include Parkinson's disease, dementia with Lewy body formation, multiple system atrophy, Hallerboden-Spatz disease, and diffuse Lewy body disease.

One of the competing theories of the aetiology of AD is that the
30 causative step(s) lies within the pathway of the intracerebral biogenesis and accumulation of the A β amyloid protein (see recent reviews by Selkoe, 2001; Beyreuther et al., 2001; Bush, 2001). However, to date no drugs or agents which target this pathway have been demonstrated to have a lasting effect on modifying the clinical expression of the disease or in preventing or ameliorating the decline in cognitive
35 function associated with neurodegenerative disorders, including Alzheimer's disease.

A further hypothesis is that AD is caused by the toxic accumulation of A β amyloid, due in part to excess binding of copper and zinc, metal ions which are

abundant in the regions most affected. Moreover, it has been suggested that when Zn^{2+} and Cu^{2+} ions interact with A β , aggregation of A β into fibrils and plaques occurs (Atwood et al., 1998); confirmed by recent data from animals deficient in synaptic Zn^{2+} (Lee et al., 2002). It has also been suggested that redox-active Cu^{2+} -A β interactions can generate H_2O_2 from O_2 (Huang et al., 1999). Both Cu^{2+} and Zn^{2+} have been shown to affect A β -lipid membrane interactions (Curtain et al., 2001).

A method of treatment of AD using iodochlorohydroxyquinoline an antibiotic [also known as clioquinol (CQ)], is disclosed and claimed in US patent Nos. 5,994,323 and 6,001,852 by P.N. Geromylatos S.A. and in US patent application No. 09/972,913 by Bush *et al.* CQ was withdrawn as an antibiotic in 1970, because of its association with an uncommon neurological syndrome, subacute myelo-optic neuropathy (SMON), which was observed only in Japan in the 1960s, in patients thought to have received the drug over long periods and probably at doses higher than those recommended at the time (Shiraki, 1975). However, recent evidence suggests that SMON was caused by an overuse-related vitamin B12 deficiency in an exceptionally vulnerable population, and therefore could be rehabilitated for study in a clinical setting (Yassin et al., 2000; Bush and Masters, 2001).

However, no *in vivo* results in animal models or in humans are provided in the Geromylatos and Bush patents. US 5,994,323 discloses a composition comprising CQ and Vitamin B12, and its use for the treatment of "diseases or disorders responsive to CQ administration while inhibiting detrimental side effects" of CQ. These diseases include AD. US 6,001,852 discloses a method of treatment of AD using CQ, preferably together with Vitamin B12. Both US 5,994,323 and US 6,001,852 suggest a dosage of 10-750 mg per day; US 5,994,323 recommends that if treatment is over a long period CQ should be given intermittently, for up to 3 weeks at a time followed by a "wash-out" period of 1-4 weeks.

In US application No. 09/972,913 CQ is exclusively referred to in terms of its ability to disaggregate A β deposits. No other mechanism of neurotoxicity is discussed. PCT/US99/05291 by General Hospital Corporation discloses the use of CQ in combination with specific copper and zinc chelators to promote dissolution of amyloid plaques and inhibition of amyloid plaque formation and/or the production of ROS by A β .

US 6,001,852 also suggests that a composition comprising CQ and Vitamin B12 could be used in the treatment of Parkinson's disease; however, in this context it is suggested that CQ acts primarily via clearing iron from the substantia nigra.

The efficacy of CQ in the treatment of AD rests upon its ability to enter the CNS and then sequester the transition metals Cu, Zn and Fe from various A β

entities thereby reducing A β toxicity and liberating it for clearance. The effectiveness of CQ is restricted by its poor aqueous solubility which limits its oral bioavailability. CQ is also known to undergo considerable conjugative metabolism and has a history of toxicity as discussed above. The fact that CQ is a bidentate metal ligand makes
 5 necessary the commitment of at least two molecules for every metal ion captured.

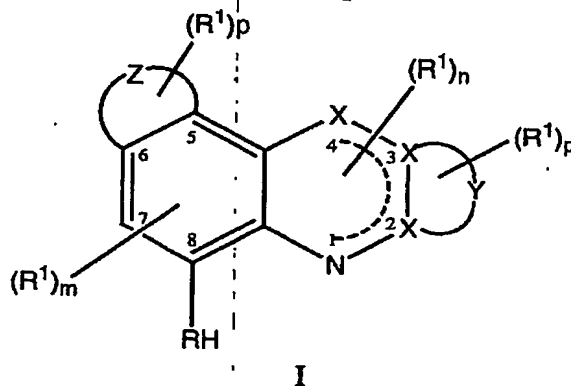
We have now developed azaquinoline derivatives which are more efficacious than CQ through the collective optimization of one or more of the following properties:

- (a) metal chelation;
- (b) aqueous solubility;
- (c) improved cell toxicity;
- (d) amyloid dispersion properties;
- (e) membrane permeability appropriate for CNS penetration; and
- (f) metabolic stability.

These derivatives include examples of therapeutics which are concentrated in the CNS through active transport, contain antioxidant activity in addition to their metal chelation properties which in some cases leads to enhanced metal chelation properties and demonstrate a prodrug strategy which masks the 8-hydroxy or 8-mercapto moiety to favour CNS penetration and make use of the known esterase
 20 activity which resides on the inner surface of the blood brain barrier (BBB).

SUMMARY OF THE INVENTION

According to the present invention there is provided a method for the
 25 treatment and/or prophylaxis of a neurological condition which comprises the administration of an effective amount of a compound of formula I:



in which

R is O or S;

R^1 is independently selected from H, optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted aryl, optionally substituted heterocyclyl, an antioxidant, a targeting moiety, CN, halo, alkylamino, alkylthio, alkylsulphonyl, alkylsulphinyl, SO_3H and OR^2 , SR^2 , NR^2R^3 , $CONR^2R^3$, $CSNR^2R^3$, $NCOR^2$, $NCSR^2$, COR^2 , CO_2R^2 , CSR^2 and $SO_2NR^2R^3$ in which R^2 and R^3 are independently selected from H, optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted aryl, optionally substituted heterocyclyl, an antioxidant or a targeting moiety;

X is independently selected from CH, CO and N;

Y and Z are independently absent or together with the ring to which they are attached form a 5- or 6-membered optionally substituted aryl or a 5- or 6-membered optionally substituted heterocyclyl;

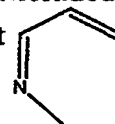
m is an integer from 1 to 3;

n is an integer from 1 to 2; and

p is an integer from 1 to 4,

salts, hydrates, solvates, derivatives, pro-drugs, tautomers and/or isomers thereof to a subject in need thereof,

with the proviso that phanquinone or tautomers thereof are excluded i.e., when R is O, R^1 at position 7 is OH, X is CH and Y is absent, then Z is not



The invention also provides use of the compound of formula I in the manufacture of a medicament for the treatment and/or prophylaxis of a neurological condition.

The invention further provides use of the compound of formula I for the treatment and/or prophylaxis of a neurological condition.

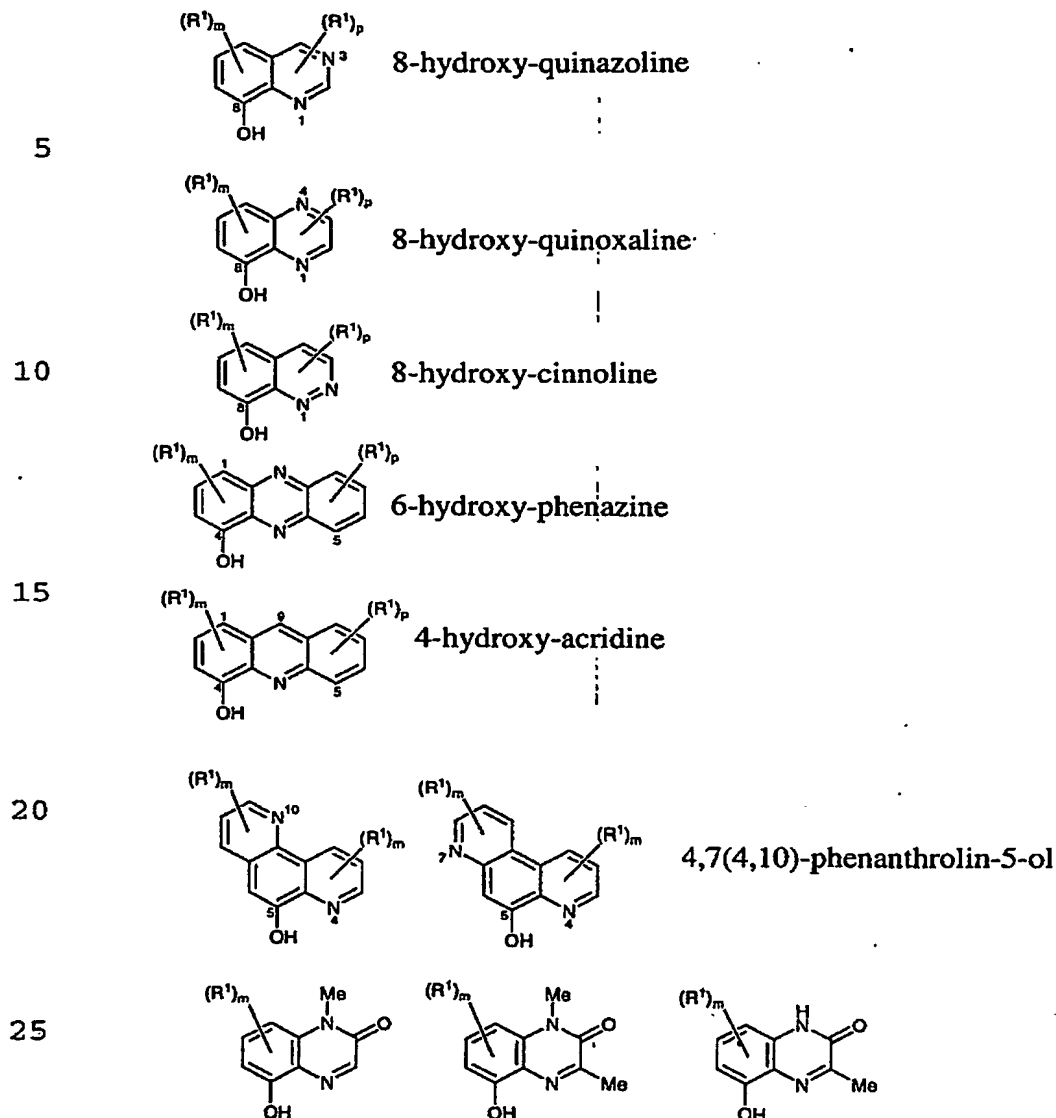
The invention still further provides use of the compound of formula I as a pharmaceutical, preferably a neurotherapeutic or neuroprotective agent, more preferably an anti-amyloidogenic agent. Preferably, the neurological condition is a neurodegenerative condition, more preferably neurodegenerative amyloidosis such as Alzheimer's disease.

Preferably, the R^1 substituent is located at positions 5 and/or 7 on formula I.

R is preferably O.

R^1 is preferably halo, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted alkyl, alkylamino, alkylthio, $CONR^2R^3$ and $NCOR^2$ in which R^2 and R^3 are as defined above.

Illustrative compounds of formula I are as follows:



in which R¹, m, n and p are as defined above.

The 8-hydroxyl or 8-mercapto group on the azaquinoline derivatives may be blocked to form a prodrug, in particular an ester prodrug. The 8-hydroxy or 8-mercapto represents a principal site of metabolism for the compound of Formula I: conjugation with glucuronic acid or sulphate gives a hydrophilic species ready to be excreted. Such conjugates probably do not pass the blood brain barrier. The ester prodrug may protect the compound of Formula I from conjugation. Esterases integral to the blood brain barrier may then release the C8-hydroxy or mercapto on passage through that barrier activating the compound for its role in the CNS.

In a further aspect, the invention provides a pharmaceutical or veterinary composition comprising the compound of formula I as defined above, together with a pharmaceutically or veterinarily acceptable carrier.

Some of the compounds of formula I are novel *per se*.

5 Accordingly, the invention provides a compound of formula II which is a compound of formula I with the provisos that when R is O and each R¹ is H or OH positioned on the rings as shown in the illustrative compounds of formula I above, then X, Y and Z are not as shown in the illustrative compounds of formula I above.

10 The compound of formula II defined above may be prepared using the processes described in detail hereinafter.

DETAILED DESCRIPTION OF THE INVENTION

For the purposes of this specification it will be clearly understood that the word "comprising" means "including but not limited to", and that the word
15 "comprises" has a corresponding meaning.

 The term "alkyl" used either alone or in compound words such as "optionally substituted alkyl" "haloalkyl" or "alkyl acyl" refers to straight chain, branched chain or cyclic hydrocarbon groups having from 1 to 10 carbon atoms, preferably 1 to 6 carbon atoms, more preferably 1 to 4 carbon atoms. Illustrative of
20 such alkyl groups are methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, neopentyl, hexyl, cyclopropyl, cyclobutyl, cyclopentyl or cyclohexyl.

 The term "alkenyl" used either alone or in compound words such as "optionally substituted alkenyl", denotes linear, branched or mono- or poly-cyclic radicals having at least one carbon-carbon double bond of 2 to 20 carbon atoms, preferably 2 to 14 carbon atoms, more preferably 2 to 6 carbon atoms. Examples of
25 alkenyl radicals include allyl, ethenyl, propenyl, butenyl, iso-butenyl, 3-methyl-2-butenyl, 1-pentenyl, cyclopentenyl, 1-methyl-cyclopentenyl, 1-hexenyl, 3-hexenyl, cyclohexenyl, 1-heptenyl, 3-heptenyl, 1-octenyl, cyclooctenyl, 1-nonenyl, 2-nonenyl, 3-nonenyl, 1-decenyl, 3-decenyl, 1,3-butadienyl, 1,4-pentadienyl, 1,3-cyclopentadienyl,
30 1,3-hexadienyl, 1,4-hexadienyl, 1,3-cyclohexadienyl, 1,4-cyclohexadienyl, 1,3-cycloheptadienyl, 1,3,5-cycloheptatrienyl, 1,3,5,7-cycloocta-tetraenyl and the like.

 The term "alkynyl" used either alone or in compound words such as "optionally substituted alkynyl" refers to straight chain or branched chain radicals having at least one carbon-carbon triple bond of 2 to 20 carbon atoms, preferably 2 to
35 14 carbon atoms, more preferably 2 to 6 carbon atoms. Examples include ethynyl, 1-propynyl, 1- and 2-butynyl, 2-methyl-2-propynyl, 2-pentynyl, 3-pentynyl, 4-pentynyl, 2-hexynyl, 3-hexynyl, 4-hexynyl, 5-hexynyl, 10-undecynyl, 4-ethyl-1-octyn-3-yl, 7-

dodecynyl, 9-dodecynyl, 10-dodecynyl, 3-methyl-1-dodecyn-3-yl, 2-tridecynyl, 11-tridecynyl, 3-tetradecynyl, 7-hexadecynyl, 3-octadecynyl and the like.

The term "heterocyclyl group" used either alone or in compound words such as "optionally substituted heterocyclyl" refers to monocyclic or polycyclic

5 heterocyclic groups containing at least one heteroatom atom selected from nitrogen, sulphur and oxygen.

Suitable heterocyclic groups include N-containing heterocyclic groups, such as, unsaturated 3 to 6-membered heteromonocyclic groups containing 1 to 4 nitrogen atoms, for example, pyrrolyl, pyrrolinyl, imidazolyl, pyrazolyl, pyridyl, 10 pyrimidinyl, pyrazinyl, pyridazinyl, triazolyl or tetrazolyl;

saturated 3 to 6-membered heteromonocyclic groups containing 1 to 4 nitrogen atoms, such as, pyrrolidinyl, imidazolidinyl, piperidino or piperazinyl;

unsaturated condensed heterocyclic groups containing 1 to 5 nitrogen atoms, such as indolyl, isoindolyl, indolizynyl, benzimidazolyl, quinolyl, isoquinolyl, 15 indazolyl, benzotriazolyl or tetrazolopyridazinyl;

unsaturated 3 to 6-membered heteromonocyclic group containing an oxygen atom, such as, pyranyl or furyl;

unsaturated 3 to 6-membered heteromonocyclic group containing 1 to 2 sulphur atoms, such as, thienyl;

20 unsaturated 3 to 6-membered heteromonocyclic group containing 1 to 2 oxygen atoms and 1 to 3 nitrogen atoms, such as, oxazolyl, isoxazolyl or oxadiazolyl;

saturated 3 to 6-membered heteromonocyclic group containing 1 to 2 oxygen atoms and 1 to 3 nitrogen atoms, such as, morpholinyl;

unsaturated condensed heterocyclic group containing 1 to 2 oxygen atoms and 1 to 3 nitrogen atoms, such as, benzoxazolyl or benzoxadiazolyl;

unsaturated 3 to 6-membered heteromonocyclic group containing 1 to 2 sulphur atoms and 1 to 3 nitrogen atoms, such as, thiazolyl or thiadiazolyl;

saturated 3 to 6-membered heteromonocyclic group containing 1 to 2 sulphur atoms and 1 to 3 nitrogen atoms, such as, thiazolidinyl; and

30 unsaturated condensed heterocyclic group containing 1 to 2 sulphur atoms and 1 to 3 nitrogen atoms, such as, benzothiazolyl or benzothiadiazolyl.

Preferably the heterocyclyl is as an unsaturated 5- or 6-membered heteromonocyclic group containing 1 to 3 nitrogen atoms such as imidazolyl, triazolyl, pyrazolyl or pyridinyl; an unsaturated condensed heterocyclic group such as quinolyl or 35 benzothiadiazolyl; an unsaturated 5-membered heteromonocyclyl group containing 1 to 2 sulphur atoms such as thiophenyl; or an unsaturated 5- or 6-membered

heteromonocyclyl group containing 1 to 2 sulphur atoms and 1 to 2 nitrogen atoms such as thiazolyl.

5 The term "aryl" used either alone or in compound words such as "optionally substituted aryl" or "aryl acyl" denotes a carbocyclic aromatic system containing one, two or three rings wherein such rings may be attached together in a pendent manner or may be fused. The term "aryl" embraces aromatic radicals such as phenyl, naphthyl, tetrahydronaphthyl, indane and biphenyl. Preferably, the aryl is phenyl.

10 The term "halo" refers to fluorine, chlorine, bromine or iodine, preferably iodine and chlorine.

The term "alkylthio" refers to radicals containing a linear or branched alkyl of 1 to 10 carbon atoms, preferably 1 to 6 carbon atoms attached to a divalent sulphur atom. Examples of alkylthio radicals include methylthio, ethylthio, propylthio, butylthio and hexylthio.

15 The term "alkylsulfinyl" refers to radicals containing a linear or branched alkyl radical, of 1 to 10 carbon atoms, preferably 1 to 6 carbon atoms attached to a divalent -S(=O)- radical. Examples include methylsulfinyl, ethylsulfinyl, butylsulfinyl and hexylsulfinyl.

20 The term "alkylsulfonyl" refers to radicals containing a linear or branched alkyl radical of 1 to 10 carbon atoms, preferably 1 to 6 carbon atoms attached to a divalent -SO₂- radical. Examples include methylsulfonyl, ethylsulfonyl and propylsulfonyl.

25 The term "optionally substituted" refers to a group which may or may not be further substituted with one or more groups selected from alkyl, alkenyl, alkynyl, aryl, aldehyde, halo, haloalkyl, haloalkenyl, haloalkynyl, haloaryl, hydroxy, alkoxy, alkenyloxy, aryloxy, benzyloxy, haloalkoxy, haloalkenyloxy, haloaryloxy, nitro, nitroalkyl, nitroalkenyl, nitroalkynyl, nitroaryl, nitroheterocyclyl, amino, alkylamino, dialkylamino, alkenylamino, alkynylamino, arylamino, diarylamino, benzylamino, dibenzylamino, acyl, alkenylacyl, alkynylacyl, arylacyl, acylamino, diacylamino, 30 acyloxy, alkylsulphonyloxy, arylsulphenyloxy, heterocyclyl, heterocycloxy, heterocyclamino, haloheterocyclyl, alkylsulphenyl, arylsulphenyl, carboalkoxy, carboaryloxy, mercapto, alkylthio, benzylthio, acylthio, phosphorus-containing groups and the like.

35 The term "antioxidant" is used herein in its broadest sense and refers to a group which has the capacity to react with a reactive oxygen species such as a hydroxyl radical in such a way as to generate a non toxic product. Examples include phenols such as 3,4,5-trimethoxyphenyl and 3,5-di-t-butyl-4-hydroxyphenyl, indole amines such

as melatonin and flavonoids. Other examples may be found the literature (Wright, 2001; Karbownik, 2001; Gilgun-Sherki, 2001).

5 The term "targeting moiety" is used herein in its broadest sense and refers to a group which will facilitate the brain delivery of the drug by way of an active transport mechanism. The targeting moiety is recognised by specific transporter enzymes integral to the blood brain barrier and these transporter enzymes then provide a mechanism for the drug to be imported into the brain. Typically such transporters are sodium dependant and their substrates contain carboxylic acids such as ascorbic acid and L-glutamate. Conjugation of the targeting moiety to the drug is enacted so as to
10 retain the acid moiety. Examples can be found in the literature (Manfredini, 2002, Sakaedu, 2001).

The term "metal chelator" is used herein in its broadest sense and refers to compounds having two or more donor atoms capable of binding to a metal atom, preferably Cu, Zn or Fe wherein at least two of the donor atoms are capable of
15 simultaneous binding to the metal atom and the resultant metal complex has a thermodynamic stability greater than or equal to that of the A β metal ion complex.

The salts of the compound of Formula I or II are preferably pharmaceutically acceptable, but it will be appreciated that non-pharmaceutically acceptable salts also fall within the scope of the present invention, since these are useful
20 as intermediates in the preparation of pharmaceutically acceptable salts. Examples of pharmaceutically acceptable salts include salts of pharmaceutically acceptable cations such as sodium, potassium, lithium, calcium, magnesium, ammonium and alkylammonium; acid addition salts of pharmaceutically acceptable inorganic acids such as hydrochloric, orthophosphoric, sulphuric, phosphoric, nitric, carbonic, boric,
25 sulfamic and hydrobromic acids; or salts of pharmaceutically acceptable organic acids such as acetic, propionic, butyric, tartaric, maleic, hydroxymaleic, fumaric, citric, lactic, mucic, gluconic, benzoic, succinic, oxalic, phenylacetic, methanesulphonic, trihalomethanesulphonic, toluenesulphonic, benzenesulphonic, salicylic, sulphanilic, aspartic, glutamic, edetic, stearic, palmitic, oleic, lauric, pantothenic, tannic, ascorbic
30 and valeric acids.

In addition, some of the compounds of the present invention may form solvates with water or common organic solvents. Such solvates are encompassed within the scope of the invention.

By "pharmaceutically acceptable derivative" is meant any
35 pharmaceutically acceptable salt, hydrate or any other compound which, upon administration to the subject, is capable of providing (directly or indirectly) a compound of Formula I or II or an active metabolite or residue thereof.

The term "pro-drug" is used herein in its broadest sense to include those compounds which are converted *in vivo* to compounds of Formula I or II. Use of the pro-drug strategy optimises the delivery of the drug to its site of action, for example, the brain. In one aspect, the term refers to the presence of a C₁₋₆ alkyl or arylester moiety which is designed to resist hydrolysis until the pro-drug has crossed the BBB, where esterases on the inner surface of the BBB act to hydrolyse the ester and liberate the C8 hydroxyl of the compounds of formula I or II. In a second aspect, the term refers to the attachment at C2 of the 8-hydroxyquinoline core of an antioxidant group, in particular the 3,4,5-trimethoxyphenyl moiety or derivatives thereof. Exposure to the prooxidative environment of the brain will then lead to hydroxylation of the 3,4,5-trimethoxyphenyl group to give a 2-hydroxy-3,4,5-trimethoxyphenyl substituent, the hydroxyl group of which acts to enhance the chelation properties of the compounds of formula I or II.

The term "tautomer" is used herein in its broadest sense to include compounds of Formula I or II which are capable of existing in a state of equilibrium between two isomeric forms. Such compounds may differ in the bond connecting two atoms or groups and the position of these atoms or groups in the compound.

The term "isomer" is used herein in its broadest sense and includes structural, geometric and stereo isomers. As the compound of Formula I or II may have one or more chiral centres, it is capable of existing in enantiomeric forms.

The compositions of the present invention comprise at least one compound of Formula I or II together with one or more pharmaceutically acceptable carriers and optionally other therapeutic agents. Each carrier, diluent, adjuvant and/or excipient must be pharmaceutically "acceptable" in the sense of being compatible with the other ingredients of the composition and not injurious to the subject. Compositions include those suitable for oral, rectal, nasal, topical (including buccal and sublingual), vaginal or parenteral (including subcutaneous, intramuscular, intravenous and intradermal) administration. The compositions may conveniently be presented in unit dosage form and may be prepared by methods well known in the art of pharmacy. Such methods include the step of bringing into association the active ingredient with the carrier which constitutes one or more accessory ingredients. In general, the compositions are prepared by uniformly and intimately bringing into association the active ingredient with liquid carriers, diluents, adjuvants and/or excipients or finely divided solid carriers or both, and then if necessary shaping the product.

The term "neurological condition" is used herein in its broadest sense and refers to conditions in which various cell types of the nervous system are degenerated and/or have been damaged as a result of neurodegenerative disorders or injuries or exposures. In particular, compounds of formula I or II can be used for the

treatment of resulting conditions, in which damage to cells of the nervous system has occurred due to surgical interventions, infections, exposure to toxic agents, tumours, nutritional deficits or metabolic disorders. In addition, compounds of the formula I or II can be used for the treatment of the sequelae of neurodegenerative disorders, such as

5 Alzheimer's disease, Parkinson's disease, multiple sclerosis, amyotrophic lateral sclerosis, epilepsy, drug abuse or drug addiction (alcohol, cocaine, heroin, amphetamine or the like), spinal cord disorders and/or injuries, dystrophy or degeneration of the neural retina (retinopathies) and peripheral neuropathies, such as diabetic neuropathy and/or the peripheral neuropathies induced by toxins

10 The term "neurodegenerative disorder" as used herein refers to an abnormality in which neuronal integrity is threatened. Neuronal integrity can be threatened when neuronal cells display decreased survival or when the neurons can no longer propagate a signal.

Neurological disorders that can be treated with the compounds of the

15 present invention include acute intermittent porphyria; adriamycin-induced cardiomyopathy; AIDS dementia and HIV-1 induced neurotoxicity; Alzheimer's disease; amyotrophic lateral sclerosis; atherosclerosis; cataract; cerebral ischaemia; cerebral palsy; cerebral tumour; chemotherapy-induced organ damage; cisplatin-induced nephrotoxicity; coronary artery bypass surgery; Creutzfeldt-Jacob disease and

20 its new variant associated with "mad cow" disease; diabetic neuropathy; Down's syndrome; drowning; epilepsy and post-traumatic epilepsy; Friedrich's ataxia; frontotemporal dementia; glaucoma; glomerulopathy; haemochromatosis; haemodialysis; haemolysis; haemolytic uraemic syndrome (Weil's disease); haemorrhagic stroke; Hallerboden-Spatz disease; heart attack and reperfusion injury;

25 Huntington's disease; Lewy body disease; intermittent claudication; ischaemic stroke; inflammatory bowel disease; macular degeneration; malaria; methanol-induced toxicity; meningitis (aseptic and tuberculous); motor neuron disease; multiple sclerosis; multiple system atrophy; myocardial ischaemia; neoplasia; Parkinson's disease; peri-natal asphyxia; Pick's disease; progressive supra-nuclear palsy; radiotherapy-induced organ

30 damage; restenosis after angioplasty; retinopathy; senile dementia; schizophrenia; sepsis; septic shock; spongiform encephalopathies; subharrachnoid haemorrhage/cerebral vasospasm; subdural haematoma; surgical trauma, including neurosurgery; thalassemia; transient ischaemic attack (TIA); traumatic brain injury (TBI); traumatic spinal injury; transplantation; vascular dementia; viral meningitis; and viral encephalitis.

35 Additionally, compounds of the present invention may also be used to potentiate the effects of other treatments, for example to potentiate the neuroprotective effects of brain derived nerve growth factor.

The invention is particularly directed to conditions which induce oxidative damage of the central nervous system, including acute and chronic neurological disorders such as traumatic brain injury, spinal cord injury, cerebral ischaemia, stroke (ischaemic and haemorrhagic), subharrachnoid haemorrhage/cerebral vasospasm, cerebral tumour, Alzheimer's disease, Creutzfeldt-Jacob disease and its new variant associated with "mad cow" disease, Huntington's disease, Parkinson's disease, Friedrich's ataxia, cataract, dementia with Lewy body formation, multiple system atrophy, Hallerboden-Spatz disease, diffuse Lewy body disease, amyotrophic lateral sclerosis, motor neuron disease and multiple sclerosis.

More particularly, the invention is directed to the treatment of neurodegenerative amyloidosis. The neurodegenerative amyloidosis may be any condition in which neurological damage results from the deposition of amyloid. The amyloid may be formed from a variety of protein or polypeptide precursors, including but not limited to A β , synuclein, huntingtin, or prion protein.

Thus the condition is preferably selected from the group consisting of sporadic or familial Alzheimer's disease, amyotrophic lateral sclerosis, motor neuron disease, cataract, Parkinson's disease, Creutzfeldt-Jacob disease and its new variant associated with "mad cow" disease, Huntington's disease, dementia with Lewy body formation, multiple system atrophy, Hallerboden-Spatz disease, and diffuse Lewy body disease.

More preferably the neurodegenerative amyloidosis is an A β -related condition, such as Alzheimer's disease or dementia associated with Down syndrome or one of several forms of autosomal dominant forms of familial Alzheimer's disease (reviewed in St George-Hyslop, 2000). Most preferably the A β -related condition is Alzheimer's disease.

In a particularly preferred embodiment of all aspects of the invention, prior to treatment the subject has moderately or severely impaired cognitive function, as assessed by the Alzheimer's Disease Assessment Scale (ADAS)-cog test, for example an ADAS-cog value of 25 or greater.

In addition to slowing or arresting the cognitive decline of a subject, the methods and compounds of the invention may also be suitable for use in the treatment or prevention of neurodegenerative conditions, or may be suitable for use in alleviating the symptoms of neurodegenerative conditions. The compounds may be able to provide at least a partial reversal of the cognitive decline experienced by patients. If administered to a subject who has been identified as having an increased risk of a predisposition to neurodegenerative conditions, or to a subject exhibiting pre-clinical manifestations of cognitive decline, such as Mild Cognitive Impairment or minimal

progressive cognitive impairment, these methods and compounds may be able to prevent or delay the onset of clinical symptoms, in addition to the effect of slowing or reducing the rate of cognitive decline.

Currently Alzheimer's disease and other dementias are usually not diagnosed until one or more warning symptoms have appeared. These symptoms constitute a syndrome known as Mild Cognitive Impairment (MCI), which was recently defined by the American Academy of Neurology, and refers to the clinical state of individuals who have memory impairment, but who are otherwise functioning well, and who do not meet clinical criteria for dementia (Petersen et al., 2001). Symptoms of MCI include:

- (1) Memory loss which affects job skills
- (2) Difficulty performing familiar tasks
- (3) Problems with language
- (4) Disorientation as to time and place (getting lost)
- (5) Poor or decreased judgement
- (6) Problems with abstract thinking
- (7) Misplacing things
- (8) Changes in mood or behaviour
- (9) Changes in personality
- (10) Loss of initiative

MCI can be detected using conventional cognitive screening tests, such as the Mini Mental Status Exam, and the Memory Impairment Screen, and neuropsychological screening batteries.

The term "subject" as used herein refers to any animal having a disease or condition which requires treatment with a pharmaceutically-active agent. The subject may be a mammal, preferably a human, or may be a domestic or companion animal. While it is particularly contemplated that the compounds of the invention are suitable for use in medical treatment of humans, it is also applicable to veterinary treatment, including treatment of companion animals such as dogs and cats, and domestic animals such as horses, ponies, donkeys, mules, llama, alpaca, pigs, cattle and sheep, or zoo animals such as primates, felids, canids, bovids, and ungulates.

Suitable mammals include members of the Orders Primates, Rodentia, Lagomorpha, Cetacea, Carnivora, Perissodactyla and Artiodactyla. Members of the Orders Perissodactyla and Artiodactyla are particularly preferred because of their similar biology and economic importance.

For example, Artiodactyla comprises approximately 150 living species distributed through nine families: pigs (Suidae), peccaries (Tayassuidae),

hippopotamuses (Hippopotamidae), camels (Camelidae), chevrotains (Tragulidae), giraffes and okapi (Giraffidae), deer (Cervidae), pronghorn (Antilocapridae), and cattle, sheep, goats and antelope (Bovidae). Many of these animals are used as feed animals in various countries. More importantly, many of the economically important animals such as goats, sheep, cattle and pigs have very similar biology and share high degrees of genomic homology.

The Order Perissodactyla comprises horses and donkeys, which are both economically important and closely related. Indeed, it is well known that horses and donkeys interbreed.

As used herein, the term "therapeutically effective amount" is meant an amount of a compound of the present invention effective to yield a desired therapeutic response, for example, to prevent or treat a neurological condition.

The specific "therapeutically effective amount" will, obviously, vary with such factors as the particular condition being treated, the physical condition of the subject, the type of subject being treated, the duration of the treatment, the nature of concurrent therapy (if any), and the specific formulations employed and the structure of the compound or its derivatives.

The compounds of the present invention may additionally be combined with other medicaments to provide an operative combination. It is intended to include any chemically compatible combination of pharmaceutically-active agents, as long as the combination does not eliminate the activity of the compound of formula I or II. It will be appreciated that the compound of the invention and the other medicament may be administered separately, sequentially or simultaneously.

Other medicaments may include, for example, where the condition is a β -amyloid related condition, particularly Alzheimer's disease, an inhibitor of the acetylcholinesterase active site, for example phenserine, galantamine, or tacrine; an antioxidant, such as Vitamin E or Vitamin C; an anti-inflammatory agent such as flurbiprofen or ibuprofen optionally modified to release nitric oxide (for example NCX-2216, produced by NicOx) or an oestrogenic agent such as 17- β -oestradiol.

Methods and pharmaceutical carriers for preparation of pharmaceutical compositions are well known in the art, as set out in textbooks such as Remington's Pharmaceutical Sciences, 20th Edition, Williams & Wilkins, Pennsylvania, USA.

As used herein, a "pharmaceutical carrier" is a pharmaceutically acceptable solvent, suspending agent or vehicle for delivering the compound of formula I or II to the subject. The carrier may be liquid or solid and is selected with the planned manner of administration in mind. Each carrier must be pharmaceutically "acceptable"

in the sense of being compatible with other ingredients of the composition and non injurious to the subject.

5 The compound of formula I or II may be administered orally, topically, or parenterally in dosage unit formulations containing conventional non-toxic pharmaceutically acceptable carriers, adjuvants, and vehicles. The term parenteral as used herein includes subcutaneous injections, aerosol for administration to lungs or nasal cavity, intravenous, intramuscular, intrathecal, intracranial, injection or infusion techniques.

10 The present invention also provides suitable topical, oral, and parenteral pharmaceutical formulations for use in the novel methods of treatment of the present invention. The compounds of the present invention may be administered orally as tablets, aqueous or oily suspensions, lozenges, troches, powders, granules, emulsions, capsules, syrups or elixirs. The composition for oral use may contain one or more agents selected from the group of sweetening agents, flavouring agents, colouring
15 agents and preserving agents in order to produce pharmaceutically elegant and palatable preparations. Suitable sweeteners include sucrose, lactose, glucose, aspartame or saccharin. Suitable disintegrating agents include corn starch, methylcellulose, polyvinylpyrrolidone, xanthan gum, bentonite, alginic acid or agar. Suitable flavouring agents include peppermint oil, oil of wintergreen, cherry, orange or raspberry
20 flavouring. Suitable preservatives include sodium benzoate, vitamin E, alphanatocopherol, ascorbic acid, methyl paraben, propyl paraben or sodium bisulphite. Suitable lubricants include magnesium stearate, stearic acid, sodium oleate, sodium chloride or talc. Suitable time delay agents include glyceryl monostearate or glyceryl distearate. The tablets contain the active ingredient in admixture with non-toxic
25 pharmaceutically acceptable excipients which are suitable for the manufacture of tablets.

These excipients may be, for example, (1) inert diluents, such as calcium carbonate, lactose, calcium phosphate or sodium phosphate; (2) granulating and disintegrating agents, such as corn starch or alginic acid; (3) binding agents, such as
30 starch, gelatin or acacia; and (4) lubricating agents, such as magnesium stearate, stearic acid or talc. These tablets may be uncoated or coated by known techniques to delay disintegration and absorption in the gastrointestinal tract and thereby provide a sustained action over a longer period. For example, a time delay material such as glyceryl monostearate or glyceryl distearate may be employed. Coating may also be
35 performed using techniques described in the U.S. Pat. Nos. 4,256,108; 4,160,452; and 4,265,874 to form osmotic therapeutic tablets for control release.

The compound of formula I or II as well as the pharmaceutically-active agent useful in the method of the invention can be administered, for *in vivo* application, parenterally by injection or by gradual perfusion over time independently or together. Administration may be intravenously, intraarterial, intraperitoneally, intramuscularly, subcutaneously, intracavity, transdermally or infusion by, for example, osmotic pump. For *in vitro* studies the agents may be added or dissolved in an appropriate biologically acceptable buffer and added to a cell or tissue.

Preparations for parenteral administration include sterile aqueous or non-aqueous solutions, suspensions, and emulsions. Examples of non-aqueous solvents are propylene glycol, polyethylene glycol, vegetable oils such as olive oil, and injectable organic esters such as ethyl oleate. Aqueous carriers include water, alcoholic/aqueous solutions, emulsions or suspensions, including saline and buffered media. Parenteral vehicles include sodium chloride solution, Ringer's dextrose, dextrose and sodium chloride, lactated Ringer's intravenous vehicles include fluid and nutrient replenishers, electrolyte replenishers (such as those based on Ringer's dextrose), and the like. Preservatives and other additives may also be present such as, for example, anti-microbials, anti-oxidants, chelating agents, growth factors and inert gases and the like.

Generally, the terms "treating", "treatment" and the like are used herein to mean affecting a subject, tissue or cell to obtain a desired pharmacologic and/or physiologic effect. The effect may be prophylactic in terms of completely or partially preventing a disease or sign or symptom thereof, and/or may be therapeutic in terms of a partial or complete cure of a disease. "Treating" as used herein covers any treatment of, or prevention of disease in a vertebrate, a mammal, particularly a human, and includes: (a) preventing the disease from occurring in a subject that may be predisposed to the disease, but has not yet been diagnosed as having it; (b) inhibiting the disease, i.e., arresting its development; or (c) relieving or ameliorating the effects of the disease, i.e., cause regression of the effects of the disease.

The invention includes various pharmaceutical compositions useful for ameliorating disease. The pharmaceutical compositions according to one embodiment of the invention are prepared by bringing a compound of formula I or II, analogues, derivatives or salts thereof, or combinations of compound of formula I or II and one or more pharmaceutically-active agents into a form suitable for administration to a subject using carriers, excipients and additives or auxiliaries. Frequently used carriers or auxiliaries include magnesium carbonate, titanium dioxide, lactose, mannitol and other sugars, talc, milk protein, gelatin, starch, vitamins, cellulose and its derivatives, animal and vegetable oils, polyethylene glycols and solvents, such as sterile water, alcohols, glycerol and polyhydric alcohols. Intravenous vehicles include fluid and nutrient

replenishers. Preservatives include antimicrobial, anti-oxidants, chelating agents and inert gases. Other pharmaceutically acceptable carriers include aqueous solutions, non-toxic excipients, including salts, preservatives, buffers and the like, as described, for instance, in Remington's Pharmaceutical Sciences, 20th ed. Williams and Wilkins
5 (2000) and The British National Formulary 43rd ed. (British Medical Association and Royal Pharmaceutical Society of Great Britain, 2002; <http://bnf.rhn.net>), the contents of which are hereby incorporated by reference. The pH and exact concentration of the various components of the pharmaceutical composition are adjusted according to routine skills in the art. See Goodman and Gilman's The Pharmacological Basis for
10 Therapeutics (7th ed., 1985).

The pharmaceutical compositions are preferably prepared and administered in dose units. Solid dose units may be tablets, capsules and suppositories. For treatment of a subject, depending on activity of the compound, manner of
administration, nature and severity of the disorder, age and body weight of the subject,
15 different daily doses can be used. Under certain circumstances, however, higher or lower daily doses may be appropriate. The administration of the daily dose can be carried out both by single administration in the form of an individual dose unit or else several smaller dose units and also by multiple administration of subdivided doses at specific intervals.

The pharmaceutical compositions according to the invention may be administered locally or systemically in a therapeutically effective dose. Amounts effective for this use will, of course, depend on the severity of the disease and the weight and general state of the subject. Typically, dosages used *in vitro* may provide useful guidance in the amounts useful for *in situ* administration of the pharmaceutical
25 composition, and animal models may be used to determine effective dosages for treatment of the cytotoxic side effects. Various considerations are described, e.g., in Langer, Science, 249: 1527, (1990). Formulations for oral use may be in the form of hard gelatin capsules wherein the active ingredient is mixed with an inert solid diluent, for example, calcium carbonate, calcium phosphate or kaolin. They may also be in the
30 form of soft gelatin capsules wherein the active ingredient is mixed with water or an oil medium, such as peanut oil, liquid paraffin or olive oil.

Aqueous suspensions normally contain the active materials in admixture with excipients suitable for the manufacture of aqueous suspension. Such excipients may be (1) suspending agent such as sodium carboxymethyl cellulose, methyl cellulose,
35 hydroxypropylmethylcellulose, sodium alginate, polyvinylpyrrolidone, gum tragacanth and gum acacia; (2) dispersing or wetting agents which may be (a) naturally occurring phosphatide such as lecithin; (b) a condensation product of an alkylene oxide with a

fatty acid, for example, polyoxyethylene stearate; (c) a condensation product of ethylene oxide with a long chain aliphatic alcohol, for example, heptadecaethylenoxycetanol; (d) a condensation product of ethylene oxide with a partial ester derived from a fatty acid and hexitol such as polyoxyethylene sorbitol monooleate, or (e) a condensation product of ethylene oxide with a partial ester derived from fatty acids and hexitol anhydrides, for example polyoxyethylene sorbitan monooleate.

The pharmaceutical compositions may be in the form of a sterile injectable aqueous or oleagenous suspension. This suspension may be formulated according to known methods using those suitable dispersing or wetting agents and suspending agents which have been mentioned above. The sterile injectable preparation may also be a sterile injectable solution or suspension in a non-toxic parenterally-acceptable diluent or solvent, for example, as a solution in 1,3-butanediol. Among the acceptable vehicles and solvents that may be employed are water, Ringer's solution, and isotonic sodium chloride solution. In addition, sterile, fixed oils are conventionally employed as a solvent or suspending medium. For this purpose, any bland fixed oil may be employed including synthetic mono- or diglycerides. In addition, fatty acids such as oleic acid find use in the preparation of injectables.

Compounds of formula I or II may also be administered in the form of liposome delivery systems, such as small unilamellar vesicles, large unilamellar vesicles, and multilamellar vesicles. Liposomes can be formed from a variety of phospholipids, such as cholesterol, stearylamine, or phosphatidylcholines.

The compounds of formula I or II may also be presented for use in the form of veterinary compositions, which may be prepared, for example, by methods that are conventional in the art. Examples of such veterinary compositions include those adapted for:

(a) oral administration, external application, for example drenches (e.g. aqueous or non-aqueous solutions or suspensions); tablets or boluses; powders, granules or pellets for admixture with feed stuffs; pastes for application to the tongue;

(b) parenteral administration for example by subcutaneous, intramuscular or intravenous injection, e.g. as a sterile solution or suspension; or (when appropriate) by intramammary injection where a suspension or solution is introduced in the udder via the teat;

(c) topical applications, e.g. as a cream, ointment or spray applied to the skin; or

(d) intravaginally, e.g. as a pessary, cream or foam.

Dosage levels of the compound of formula I or II of the present invention are of the order of about 0.5 mg to about 20 mg per kilogram body weight,

with a preferred dosage range between about 0.5 mg to about 10 mg per kilogram body weight per day (from about 0.5 gms to about 3 gms per patient per day). The amount of active ingredient that may be combined with the carrier materials to produce a single dosage will vary depending upon the host treated and the particular mode of administration. For example, a formulation intended for oral administration to humans may contain about 5 mg to 1g of an active compound with an appropriate and convenient amount of carrier material which may vary from about 5 to 95 percent of the total composition. Dosage unit forms will generally contain between from about 5 mg to 500 mg of active ingredient.

Optionally the compounds of the invention are administered in a divided dose schedule, such that there are at least two administrations in total in the schedule. Administrations are given preferably at least every two hours for up to four hours or longer; for example the compound may be administered every hour or every half hour. In one preferred embodiment, the divided-dose regimen comprises a second administration of the compound of the invention after an interval from the first administration sufficiently long that the level of active compound in the blood has decreased to approximately from 5-30% of the maximum plasma level reached after the first administration, so as to maintain an effective content of active agent in the blood. Optionally one or more subsequent administrations may be given at a corresponding interval from each preceding administration, preferably when the plasma level has decreased to approximately from 10-50% of the immediately-preceding maximum.

It will be understood, however, that the specific dose level for any particular patient will depend upon a variety of factors including the activity of the specific compound employed, the age, body weight, general health, sex, diet, time of administration, route of administration, rate of excretion, drug combination and the severity of the particular disease undergoing therapy.

EXAMPLES

The invention will now be described in detail by way of reference only to the following non-limiting examples.

General

The following compounds were prepared by the methods described in the literature: 8-methoxy-4(3*H*)-quinazolinone (3),⁵ 8-hydroxy-cinnoline (C1),² 5,8-dihydroxy-quinoxaline (B39),¹ 4,5-dihydroxy-phenazine (F5),³ 4,8-dihydroxy-

phenazine (F2),³ 4-hydroxy-acridine (E1),⁴ and 5-hydroxy-3-methyl-2(1H)-quinoxalinone (B12)¹⁹. The following compounds were sourced commercially: 8-Hydroxy-quinazoline (A1), 4-hydroxy-phenazine (F1), 4,10-phenanthroline-5-ol (D2), 4,7-phenanthroline-5,6-diol (D3) and 8-hydroxy-2-methyl-4(3H)-quinazolinone (1).

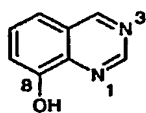
5 Amines: ethylamine, histamine, 2-(2-aminoethyl)pyridine, 2-(2-methylaminoethyl)pyridine; aldehydes: 4-imidazolecarboxaldehyde, 2-thiazolecarboxaldehyde and 2-pyridinecarboxaldehyde, azoles: pyrazole, imidazole, methylimidazole and 1H-1,2,3-triazole, boronic acids: phenylboronic acid, 2-(trifluoromethyl)phenylboronic acid, 2-methoxyphenylboronic acid, *o*-tolylboronic acid, 2-fluorophenylboronic acid, 3-methoxyphenylboronic acid, 4-10 methoxyphenylboronic acid, *m*-tolylboronic acid, 3,5-difluorophenylboronic acid, 2,4-difluorophenylboronic acid, 3-thiopheneboronic acid, 3-fluorophenylboronic acid and 4-fluorophenylboronic acid; and organozinc reagents: 2-pyridylzinc bromide, 2-(methylthio)phenylzinc iodide, 2-(ethoxycarbonyl)phenylzinc iodide and 6-15 methylpyridylzinc bromide (0.5 M solution in THF) were commercially available (Aldrich). 3-Pyridylboronic acid was purchased from Frontier Scientific. 2-Aminomethylthiazole was prepared according to the literature.¹³ Solvents were analytical grade and used as supplied. THF was distilled from sodium and benzophenone under argon. ¹H NMR spectra (δ , relative to TMS) were recorded on a 20 Varian Unity 300 spectrometer unless otherwise indicated; *J*-Values are given in hertz. Mass spectral data were recorded on a Micromass Quattro II mass spectrometer.

The synthesis of derivatives of 6 classes of compounds: 8-hydroxy-quinazoline, 8-hydroxy-quinoxaline, 8-hydroxy-cinnoline, 4,7(4,10)-phenanthroline-5-ol, 4-hydroxy-acridine and 6-hydroxy-phenazine, is described in Part A, B, C, D, E and 25 F, respectively.

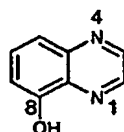
PART A: SYNTHESIS OF 8-HYDROXY-QUINAZOLINE DERIVATIVES

A2 – A138

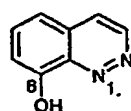
30 The preparation of a series of 8-hydroxy-quinazoline derivatives is summarized in Charts A1 – A5.



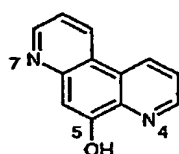
Class A



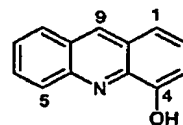
Class B



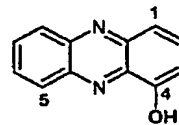
Class C



Class D

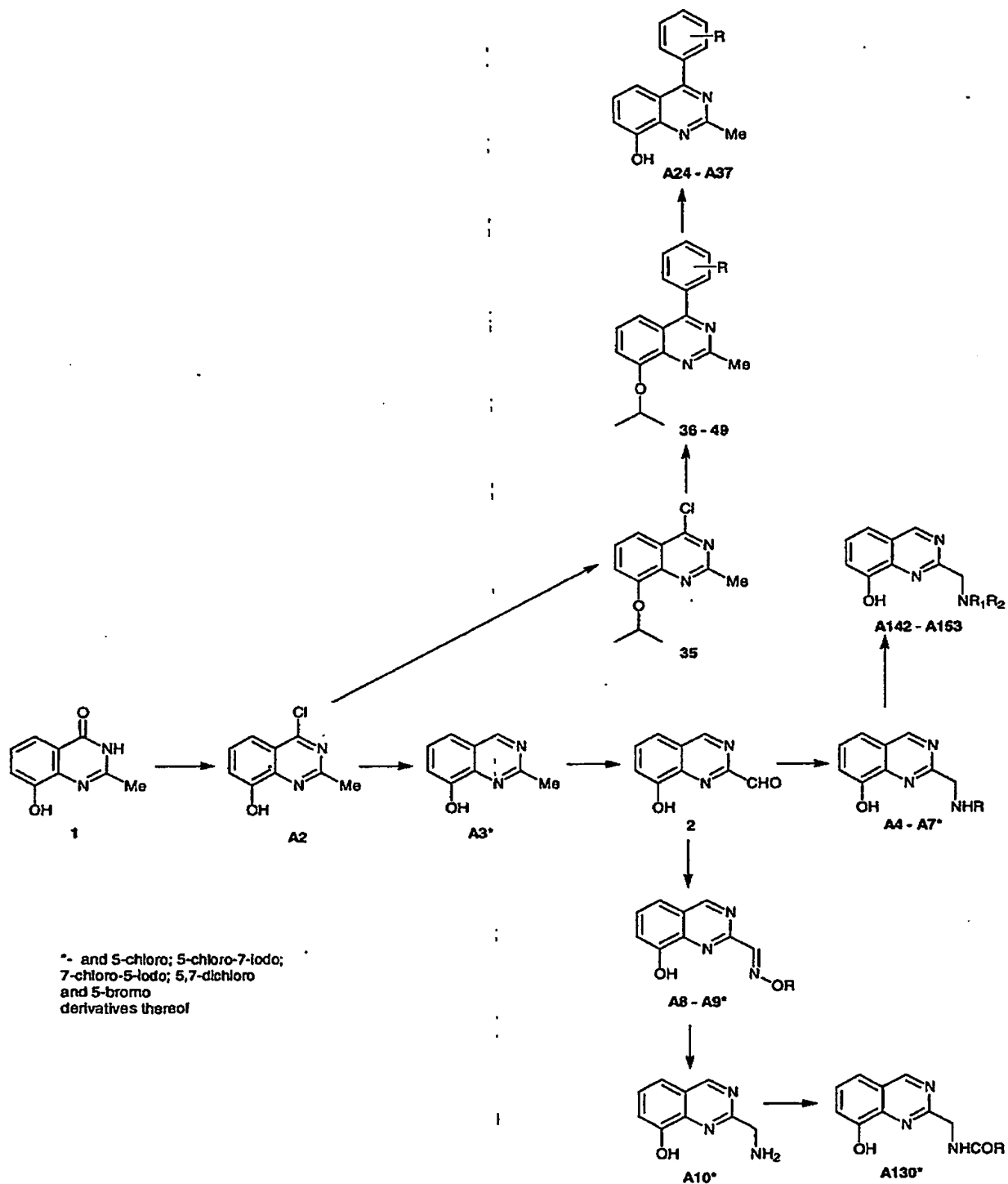


Class E

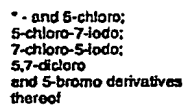


Class F

CHART A1



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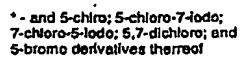


CHART A4

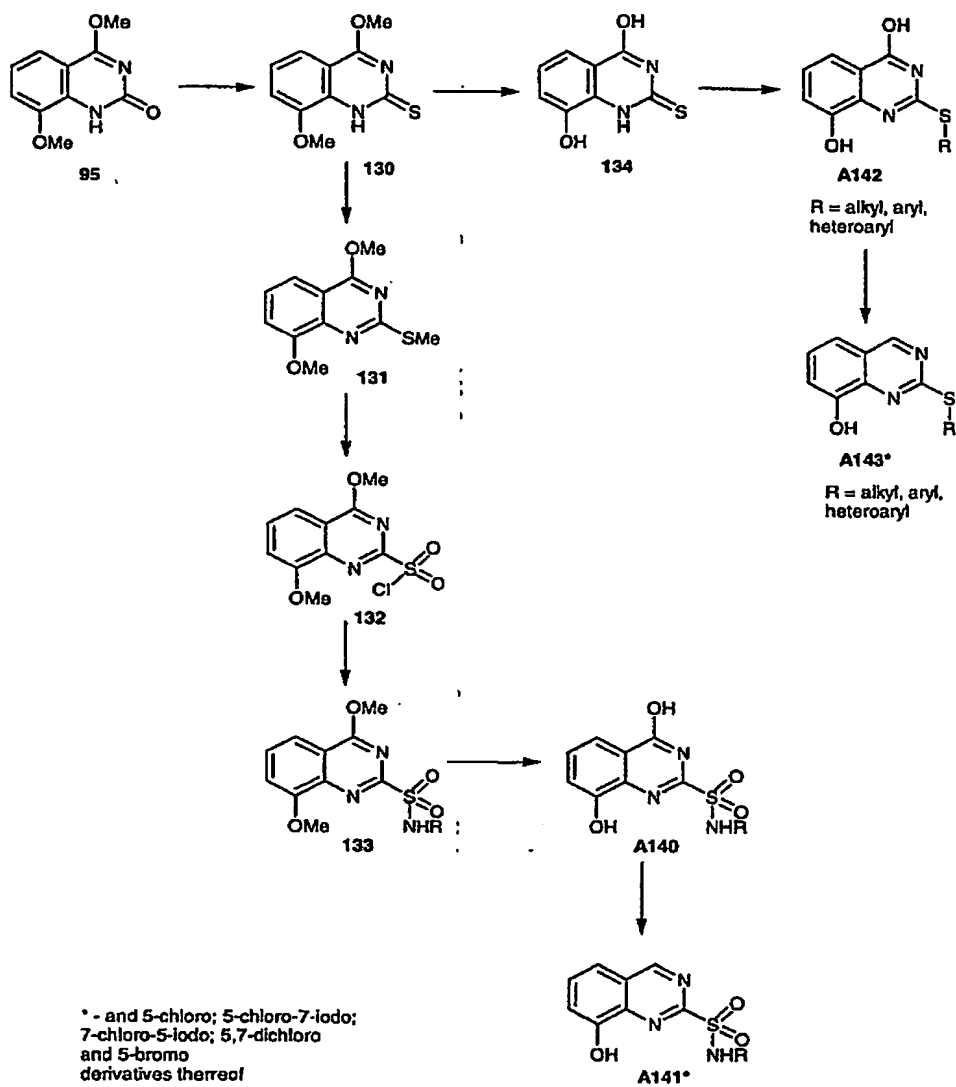
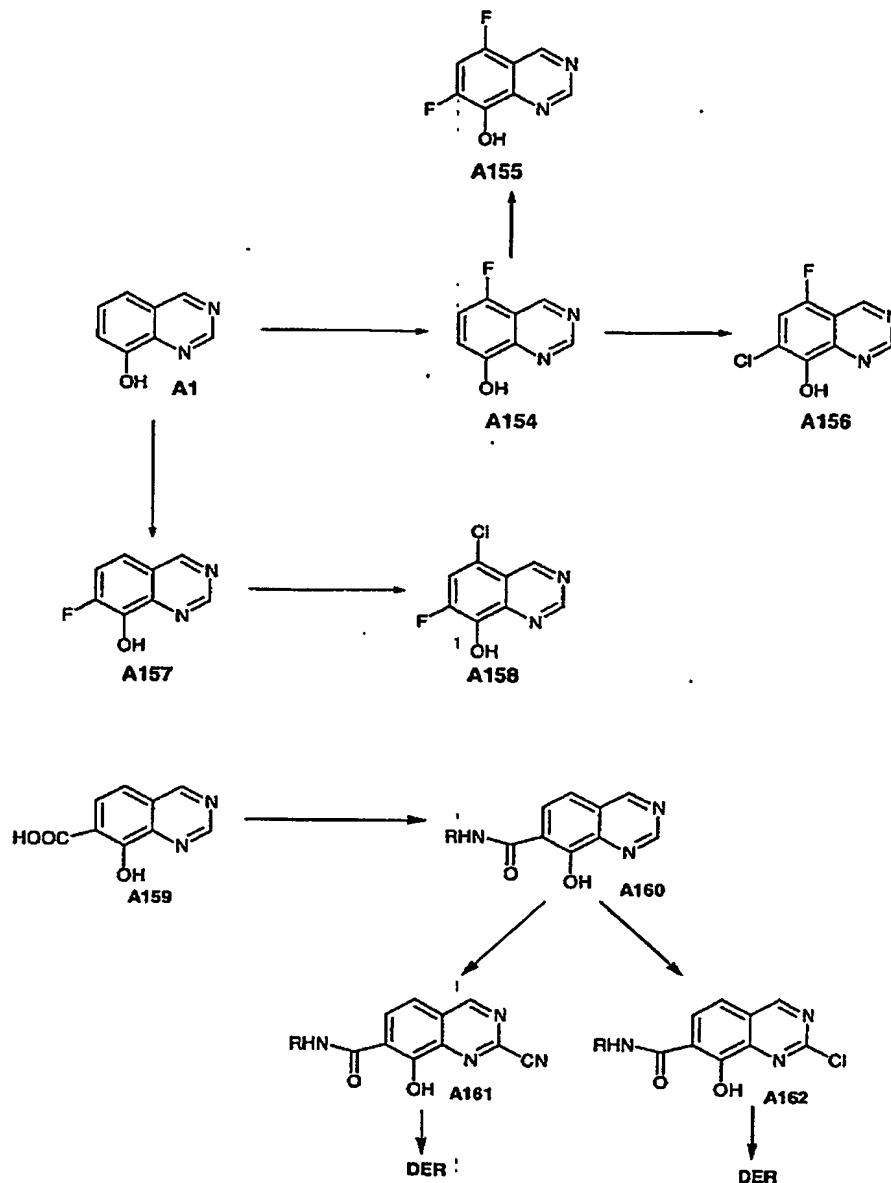
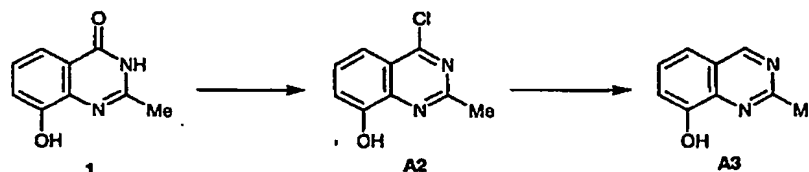


CHART A5



DER: further derivatives via substitution at the 2- (chloro/cyano) position to a subject in need thereof, as shown in CHARTS A1 - A4. 5-Position was also halogen-substituted (as shown on this CHART)

Example 1 Preparation of 4-Chloro-8-hydroxy-2-methyl-quinazoline (A2) (Scheme A1)



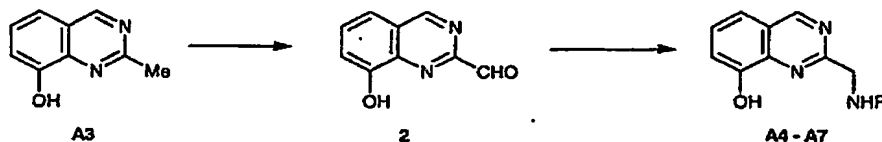
Scheme A1

5
8-Hydroxy-2-methyl-4(3H)-quinazolinone (0.01 mol) and phosphorus
oxychloride (10 mL) were heated under reflux for 30 min. The excess phosphorus
oxychloride was removed under reduce pressure and the residue was added to a mixture
of ice (50 g) and water (50ml), and the pH adjusted to 6 (aqueous ammonia). 4-Chloro-
8-hydroxy-2-methyl-quinazoline (A2) was isolated via filtration.

Preparation of 8-Hydroxy-2-methyl-quinazoline (A3) (Scheme A1)

15
4-Chloro-8-hydroxy-2-methyl-quinazoline (0.01 mol) was treated with
hydriodic acid (100 mL; freshly distilled from red phosphorus) according to the method
described⁶ in the literature. This provided 8-hydroxy-2-methyl-quinazoline (A3) as a
solid.

Example 2 Preparation of 8-Hydroxy-quinazoline-2-carboxaldehyde (2) (Scheme A2)



Scheme A2

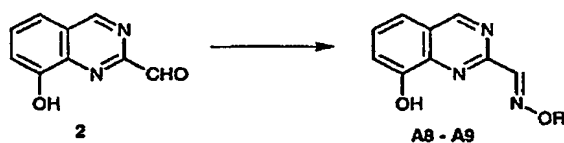
20
A solution of 8-hydroxy-2-methyl-quinazoline (A3) (5 mmol) in dioxane
(10 mL) was added dropwise over 3 h into a stirred mixture of SeO₂ (8.8 mmol) in
dioxane (30 mL) at 50°C. The resulting mixture was then heated at 80°C for 16 h,
allowed to cool, and the solids filtered off. The filtrate was concentrated and purified
via column chromatography on silica (dichloromethane/MeOH, 40:1). This afforded 8-
hydroxy-quinazoline-2-carboxaldehyde (2) as a solid.

Example 3 Preparation of 2-[Alkylamino-methyl]-8-hydroxy-quinoxaline (A4 - A7) (Scheme A2)

Sodium triacetoxyborohydride (1 mmol) was added to a stirred solution of 8-hydroxy-quinoxaline-2-carboxaldehyde (1 mmol) and ethylamine (1 mmol) in dichloromethane (10 mL). The mixture was left to stir at RT for 16 h, neutralized (aqueous NaHCO₃), and concentrated. The residue, after column chromatography on silica, afforded A4.

In a similar fashion, reductive amination of 2 with amines: histamine gave A5, 2-(2-aminoethyl)pyridine gave A6, 2-(2-methylaminoethyl)pyridine gave A7.

Example 4 Preparation of 8-hydroxy-quinazoline-2-carboxaldehyde Oximes (A8 - A9) (Scheme A3)

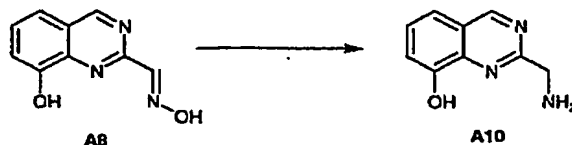


Scheme A3

A mixture of 8-hydroxy-quinazoline-2-carboxaldehyde (2) (1 mmol), NaOAc (2 mmol), hydroxylamine hydrochloride (1.5 mmol) and water (10 mL) was heated at 100 °C for 15 min. The precipitate was isolated by filtration. This provided 8-hydroxy-quinazoline-2-carboxaldehyde oxime (A8) as a solid.

The reaction, repeated using 2 with methoxylamine hydrochloride in pyridine, gave A9.

Example 5 Preparation of 8-Hydroxy-2-methylamino-quinazoline (A10) (Scheme A4)

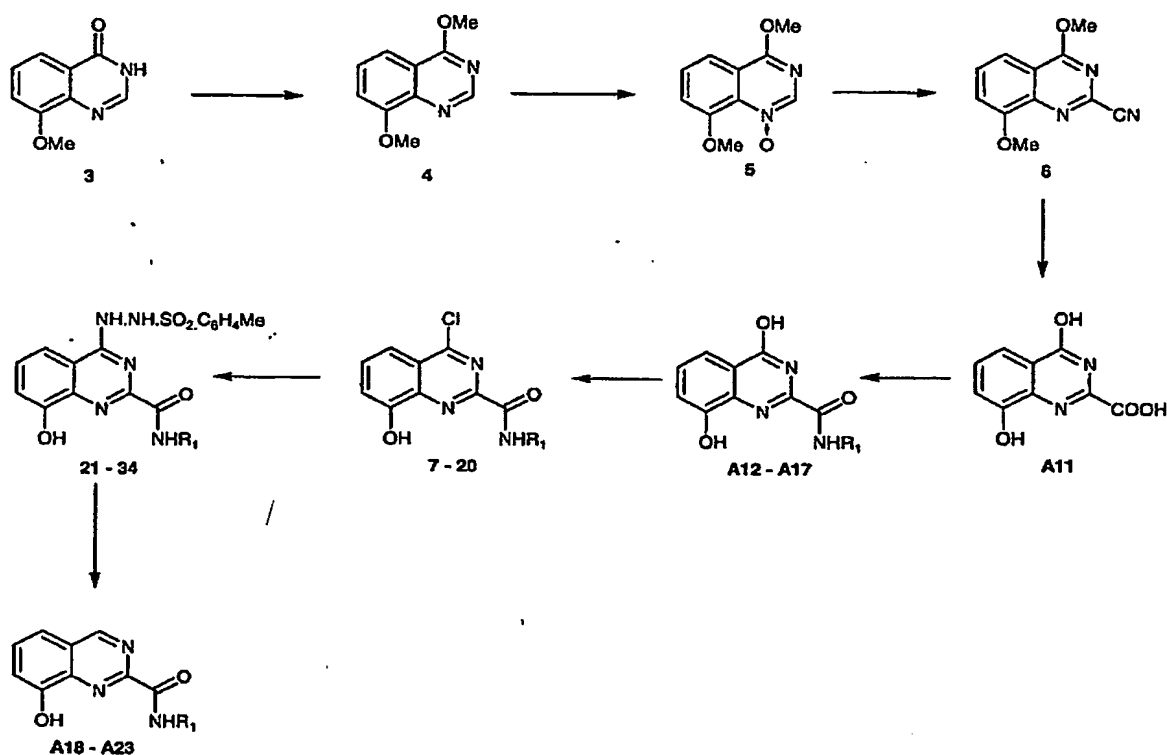


Scheme A4

A solution of the 2-carboxylic acid oxime (1 mmol) in MeOH (20 mL) was treated under hydrogenolysis conditions (atmospheric H₂, 10% Pd/C) for 16 h. The solids were filtered off and the filtrate concentrated to provide 8-hydroxy-2-methylamino-quinazoline (A10).

5

Example 6 Preparation of 4,8-Dimethoxy-quinazoline-2-carboxylic acid (A11) (Scheme A5)



Scheme A5

10

To a stirring mixture of 8-methoxy-4(3H)-quinazolinone⁵(3) (0.05 mol) and THF (100 mL) was added iodomethane (0.1 mol), tetrabutylammonium bromide (100 mg) and aqueous NaOH (prepared from 7.55 g of NaOH in 20 mL H₂O). After 16 h at 40 °C, the mixture was concentrated and the remaining residue partitioned between H₂O and dichloromethane (1:1, 200 mL). The organic layer was washed with brine, dried and concentrated. Column purification gave 4,8-dimethoxy-quinazoline (4).

15

To a stirred solution of 4 (40 mmol) in CHCl₃ (200 mL) at 0 °C was added *m*-chloroperbenzoic acid (44 mmol) portionwise over 10 min. After a further 30 min at 0 °C, the mixture was allowed to warm to RT over 30 min and then concentrated to dryness. To the remaining residue was added ethyl acetate and 1 N NaHCO₃ (1:1,

20

200 mL); the layers were separated and the organic layer was dried (Na_2SO_4), and concentrated. This provided the *N*-oxide 5.

5 A mixture of 5 (30 mmol), benzene (80 mL) and dimethyl sulphate (35 mmol) was stirred under reflux for 16 h, allowed to cool, and concentrated *in vacuo*. To the remaining residue in H_2O (100 mL) at 0 °C was added NaCN (90 mmol). After 3 h, the reaction mixture was neutralised (HOAc) and extracted with dichloromethane, the extracts combined and dried. Solvent removal gave the 2-cyano-compound 6.

10 A mixture of 6 (20 mmol) and NaOH (40 mmol) in H_2O (20 mL) was heated at 100 °C for 4 h, and cooled. The pH of the solution was adjusted to 4 (glacial HOAc) and the mixture extracted with ethyl acetate (50 mL x 4). The combined extracts were dried and the volatiles removed. This provided 4,8-dimethoxy-quinazoline-2-carboxylic acid as a solid. Subsequent de-*O*-methylation with BBr_3 gave 4,8-dihydroxy-quinazoline-2-carboxylic acid (A11).

15 **Example 7 Preparation of 4,8-dihydroxy-quinazoline-2-carboxylic acid amides (A12 - A17) (Scheme A5)**

20 1,3-Dicyclohexylcarbodiimide (1 mmol) was added to a stirred solution of 1-hydroxybenzotriazole hydrate (1 mmol) and 4,8-dihydroxy-quinazoline-2-carboxylic acid (A11) (1 mmol) in DMF and dichloromethane (1:1, 10 mL). After 30 min, histamine (1 mmol) was added and the mixture stirred at RT for a further 16 h. The volatiles were then removed *in vacuo* and the remaining residue gave, after purification by column chromatography on silica (ethyl acetate/*i*-PrOH/2 N NH_4OH , 6:2:1), 4,8-dihydroxy-quinazoline-2-carboxylic acid[2-(1*H*-imidazol-4-yl)-ethyl]-amide (A12).

25 The above reaction was repeated using amines with A11: 2-(2-aminoethyl)pyridine gave A13, 2-(aminomethyl)pyridine gave A14, 2-aminothiazole gave A15, 2-aminophenol gave A16, 1,2-phenylenediamine gave A17.

30 **Example 8 Preparation of 8-Hydroxy-quinazoline-2-carboxylic acid amides (A18 - A23) (Scheme 5)**

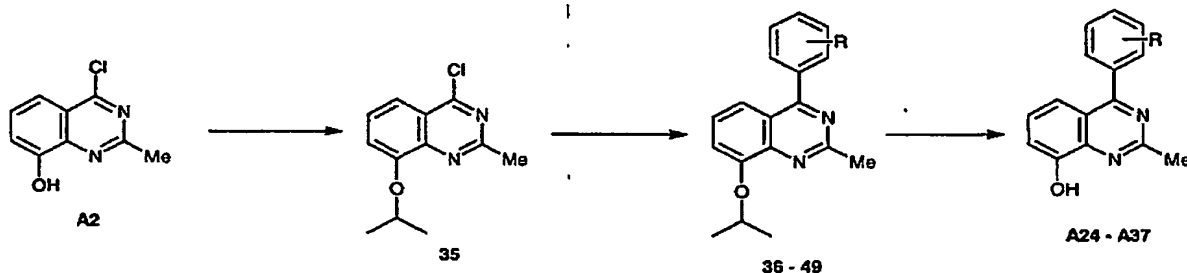
35 The 4-hydroxy-compound (A12) (1.2 mmol) and phosphorus oxychloride (4 mL) were heated under reflux for 15 min and allowed to cool. Ice (50 g) was added and the mixture basified with aqueous ammonia. The mixture was extracted with dichloromethane (20 mL x 3), the extracts combined, dried, and concentrated. This provided the corresponding 4-chloro-compound 7.

To a stirred mixture of *p*-toluenesulfonylhydrazide (Aldrich, 2 mmol) in CHCl₃ (10 mL) at 50 °C was added portionwise over 10 min, the 4-chloro-compound **7** (1 mmol). After 16 h, the solid was isolated via filtration, washed with H₂O, and dried. This provided the 4-*N'*-(*p*-toluenesulfonylhydrazino)-compound **21**.

The 4-*N'*-(*p*-toluenesulfonylhydrazino)-compound **21** (0.8 mmol) was added to Na₂CO₃ (10 mmol) and H₂O (10 mL) at 95 °C, and the mixture heated under reflux for 15 min, cooled, filtered, and the filtrate extracted with CHCl₃. The extracts were combined and dried. Subsequent removal of volatiles afforded the 8-hydroxy-quinazoline-2-carboxylic acid amide (**A18**).

In a similar fashion, the remaining 4(3*H*)-quinazolinones **A13** – **A17** were converted into 8-hydroxy-quinazoline-2-carboxylic acid amides (**A19** – **A23**).

Example 9 Preparation of 4-Aryl(or heterocyclic)-8-hydroxy-2-methyl-quinoxaline (A24– A37) (Scheme A6)



Scheme A6

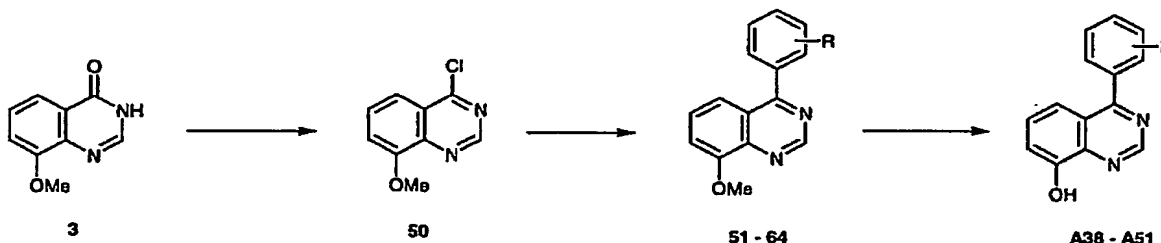
2-Bromopropane (9 mmol) was added to a stirred mixture of 4-chloro-8-hydroxy-2-methyl-quinazoline (**A2**) (6 mmol), K₂CO₃ (24 mmol) and DMSO (20 mL). After 16 h at RT, saturated NH₄Cl (20 mL) was added and the mixture extracted with dichloromethane (20 mL x 3). The extracts were combined and concentrated. Diethyl ether (100 mL) was added to the residue and the resulting mixture washed successively with 2 N NaOH, H₂O and brine, and dried (Na₂SO₄). Solvent removal afforded 4-chloro-8-isopropoxy-2-methyl-quinazoline (**35**) as a solid.

To a stirred mixture of 4-chloro-8-isopropoxy-2-methyl-quinazoline (**35**) (0.58 mmol), phenylboronic acid (0.62 mmol), 2 N Na₂CO₃ (7.2 mL), EtOH (1.2 mL) and benzene (6 mL) was added, under a blanket of argon, Pd(PPh₃)₄ (20 mg). The mixture was stirred under reflux for 16 h, cooled, and concentrated. Subsequent column chromatography (ethyl acetate/hexane) provided 8-isopropoxy-2-methyl-4-phenyl-quinazoline (**36**) as a solid.

To a stirred solution of 8-isopropoxy-2-methyl-4-phenyl-quinazoline (36) (0.34 mmol) in dichloromethane (2 mL) at -78°C was added BCl_3 (1.36 mL of a 1 M solution in dichloromethane, 1.36 mmol). The reaction mixture was allowed to warm to RT (over 2 h) and stirred for a further 2 h. MeOH (5 mL) was added and the mixture was concentrated to dryness. The process was repeated four times. Further washing of the remaining residue with diethyl ether (2 mL x 3) provided 8-hydroxy-2-methyl-4-phenyl-quinazoline (A24).

In a similar fashion, treatment of 4-chloro-8-isopropoxy-2-methyl-quinazoline (35) with boronic acids: 2-(trifluoromethyl)phenylboronic acid, 2-methoxyphenylboronic acid, *o*-tolylboronic acid, 2-fluorophenylboronic acid, 3-methoxyphenylboronic acid, 4-methoxyphenylboronic acid, *m*-tolylboronic acid, 3,5-difluorophenylboronic acid, 2,4-difluorophenylboronic acid, 3-thiopheneboronic acid, 3-fluorophenylboronic acid, 4-fluorophenylboronic acid and 3-pyridylboronic acid; and isopropoxy cleavage with BCl_3 gave 4-aryl(or heterocyclic)-8-hydroxy-2-methyl-quinazolines (A25 – A37).

Preparation of 4-Aryl(or heterocyclic)-8-hydroxy-quinazoline (A38 – A51) (Scheme A7)

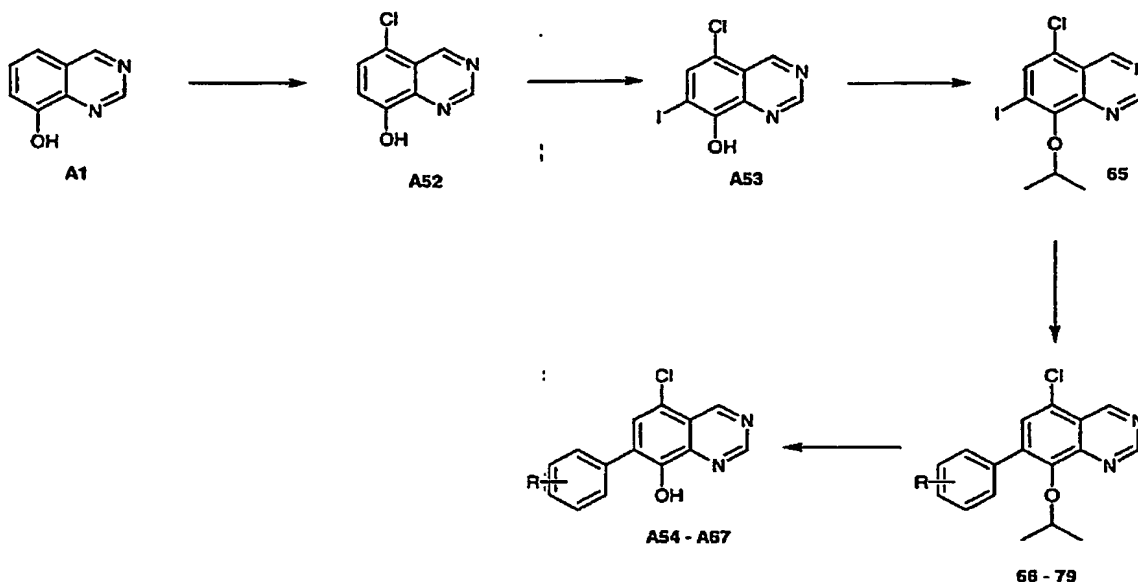


Scheme A7

Using the procedure as previously described in Example 9, 8-methoxy-4(3*H*)-quinazolinone⁵ (3) (10 mmol) and phosphorus oxychloride provided 4-chloro-8-methoxy-quinazoline 50. Treatment of the 4-chloride 50 with phenylboronic acid as described in Example 9 gave, after de-*O*-methylation with BBr_3 , the 4-phenyl derivative A38. The coupling reaction of 50 was repeated using a range of boronic acids: 2-(trifluoromethyl)phenylboronic acid, 2-methoxyphenylboronic acid, *o*-tolylboronic acid, 2-fluorophenylboronic acid, 3-methoxyphenylboronic acid, 4-methoxyphenylboronic acid, *m*-tolylboronic acid, 3,5-difluorophenylboronic acid, 2,4-difluorophenylboronic acid, 3-thiopheneboronic acid, 3-fluorophenylboronic acid, 4-fluorophenylboronic acid and 3-pyridylboronic acid gave 51 - 64; and subsequent

cleavage of the methyl ether with BBr_3 gave the 4-aryl(or heterocyclic)-8-hydroxy-quinazolines A39 – A51.

Example 10 Preparation of 5-Chloro-8-hydroxy-quinazoline (A52) (Scheme A8)



Scheme A8

Chlorine (12 mmol) was added into a stirred solution of 8-hydroxy-quinazoline (A1) (10 mmol) in 93% H_2SO_4 following a previously published⁷ procedure. After 3 h, ice (100 g) and H_2O (100 mL) was added, the mixture basified with aqueous ammonia, extracted with dichloromethane, and the extracts dried. Solvent removal gave the 5-chloro-compound A52.

Example 11 Preparation of 5-Chloro-8-hydroxy-7-iodo-quinazoline (A53) (Scheme A8)

5-Chloro-8-hydroxy-quinazoline (10 mmol) was added to a stirred solution of ICl (12 mmol) in concentrated HCl (10 mL).⁸ After 5 min, the precipitate was isolated via filtration, washed successively with H_2O , saturated sodium thiosulfate and H_2O , and dried. This provided 5-chloro-8-hydroxy-7-iodo-quinazoline (A53) as a solid.

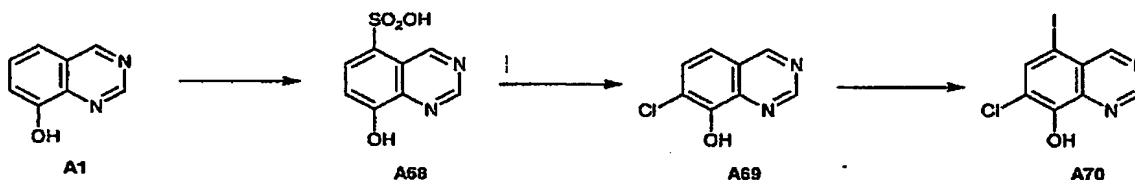
Preparation of 7-Aryl(or heterocyclic)-5-chloro-8-hydroxy-quinazolines (A54 – A67) (Scheme A8)

The 8-hydroxy-compound A53 was converted into the corresponding isopropyl ether 65 using 2-bromopropane according to the method as described in

5 Example 9.

Treatment of 5-chloro-7-iodo-8-isopropoxy-quinazoline (65), according to the method described in Example 9, with a range of boronic acids: phenylboronic acid, 2-(trifluoromethyl)phenylboronic acid, 2-methoxyphenylboronic acid, *o*-tolylboronic acid, 2-fluorophenylboronic acid, 3-methoxyphenylboronic acid, 4-methoxyphenylboronic acid, *m*-tolylboronic acid, 3,5-difluorophenylboronic acid, 2,4-difluorophenylboronic acid, 3-thiopheneboronic acid, 3-fluorophenylboronic acid, 4-fluorophenylboronic acid and 3-pyridylboronic acid gave 66 - 79; and subsequent cleavage of the 8-isopropoxy group with BCl₃, gave A54 - A79.

15 **Example 12 Preparation of 7-Chloro-8-hydroxy-quinazoline (A69) (Scheme A9)**



Scheme A9

20 8-Hydroxy-quinazoline (A1) (0.1 mol) and concentrated sulfuric acid (50 mL) were heated at 100 °C for 5 h, and allowed to cool. The solution was then carefully added to 300 mL of cold H₂O. The resulting precipitate was isolated via filtration, washed with H₂O, and dried. This provided the 5-sulfonic acid A68 as a solid.

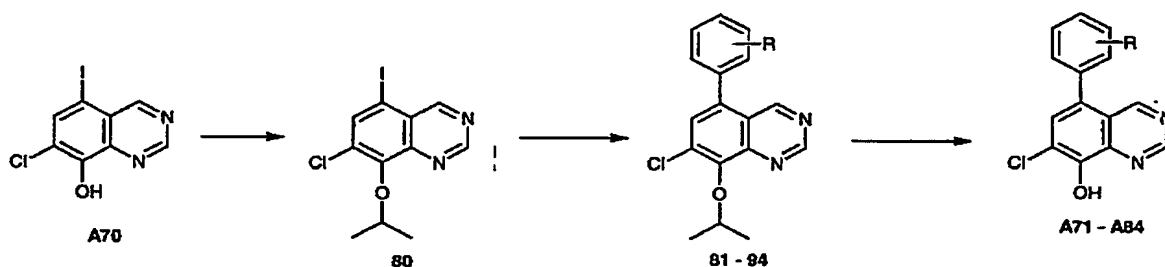
To a stirred mixture of 5-sulfonic acid A68 (0.09 mol) and H₂O (225 mL) was added KOH (0.25 mol) and NaOCl (230 mL of a solution containing 10-13% available chlorine).⁹ After 1.5 h at RT, the solution was passed through a column of Amberlite IR-120(H⁺) resin. The effluent concentrated to 20 mL. Acetone (20 mL) was then added and the precipitate isolated by filtration. Subsequent washing with acetone and drying gave 7-chloro-8-hydroxy-quinazoline (A69).

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Example 13 Preparation of 7-Chloro-8-hydroxy-5-iodo-quinazoline (A70)
(Scheme A9)

To a solution of 7-chloro-8-hydroxy-quinazoline (A69) (0.1 mol) and potassium acetate (0.15 mol) in MeOH and H₂O (19:1, 250 mL) was added, over 30 min, a solution of iodine (0.095 mol) in MeOH and H₂O (19:1, 350 mL). The mixture was then heated under reflux for 15 min, cooled and H₂O (400 mL) was added. The precipitate was isolated by filtration, washed with saturated sodium thiosulfate and H₂O, and dried. This provided 7-chloro-8-hydroxy-5-iodo-quinazoline (A70) as a solid.

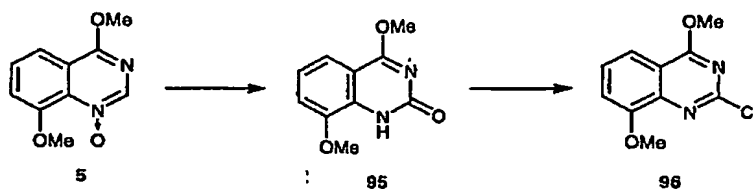
Preparation of 4-Aryl(or heterocyclic)-7-chloro-8-hydroxy-quinazolines (A71 – A84) (Scheme A10)



Scheme A10

According to the method described in Example 9, treatment of 5-chloro-7-iodo-8-isopropoxy-quinazoline (80), with a range of boronic acids: phenylboronic acid, 2-(trifluoromethyl)phenylboronic acid, 2-methoxyphenylboronic acid, *o*-tolylboronic acid, 2-fluorophenylboronic acid, 3-methoxyphenylboronic acid, 4-methoxyphenylboronic acid, *m*-tolylboronic acid, 3,5-difluorophenylboronic acid, 2,4-difluorophenylboronic acid, 3-thiopheneboronic acid, 3-fluorophenylboronic acid, 4-fluorophenylboronic acid and 3-pyridylboronic acid gave 81 - 94; and subsequent cleavage of the 8-isopropoxy group with BCl₃, gave A71 – A84.

Example 14 Preparation of 2-Chloro-4,8-dimethoxy-quinoxaline (96)
(Scheme A11)

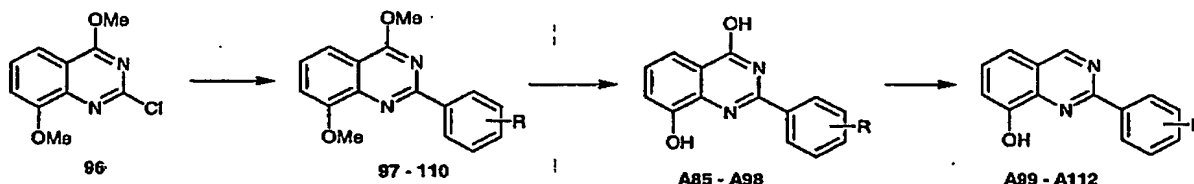


Scheme A11

Ac₂O (6 mL) was added to a stirred mixture of the *N*-oxide **5** (10 mmol) and dichloromethane (10 mL). The solvent was then removed *in vacuo* and the resulting solution heated under reflux for 1 h, cooled, and concentrated. The remaining residue was washed with diethyl ether (10 mL x 2). This provided 4,8-dimethoxy-2(1*H*)-quinazolinone (**95**) as a solid.

Treatment of 4,8-dimethoxy-2(1*H*)-quinazolinone (**95**) (5 mmol) with phosphorus oxychloride (20 mL) and standard workup according to Example 8 provided the 2-chloride **96**.

Preparation of 2-Aryl(or heterocyclic)-4,8-dihydroxy-quinazolines (A85 – A98)
(Scheme A12)



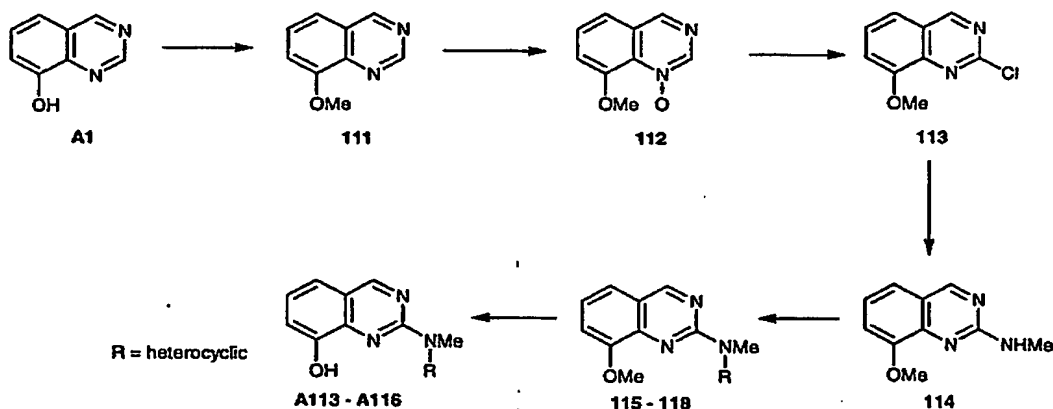
Scheme A12

According to the procedure previously described in Example 9, coupling of the 2-chloride **96** (0.1 mmol) with boronic acids: phenylboronic acid, 2-(trifluoromethyl)phenylboronic acid, 2-methoxyphenylboronic acid, *o*-tolylboronic acid, 2-fluorophenylboronic acid, 3-methoxyphenylboronic acid, 4-methoxyphenylboronic acid, *m*-tolylboronic acid, 3,5-difluorophenylboronic acid, 2,4-difluorophenylboronic acid, 3-thiophenylboronic acid, 3-fluorophenylboronic acid, 4-fluorophenylboronic acid and 3-pyridylboronic acid gave **97 - 110**; and subsequent cleavage of the 8-*O*-methyl ether with BBr₃, gave **A85 – A98**.

Preparation of 2-Aryl(or heterocyclic)-8-hydroxy-quinazoline (Scheme A12)

Sequential treatment of the 4-hydroxy-compounds A85 – A98 with phosphorus oxychloride, *p*-toluenesulfonylhydrazide and Na₂CO₃, according to the method previously described in Example 6, provided the 2-aryl(or heterocyclic)-8-hydroxy-quinazolines A99 – A112.

Preparation of 2-Chloro-8-methoxy-quinazoline (113) (Scheme A13)



Scheme A13

According to the procedure previously described in Example 6, 8-hydroxy-quinazoline (A1) (20 mmol) and iodomethane gave 8-methoxy-quinazoline (111). 8-Methoxy-quinazoline (111) (10 mmol) was then treated with *m*-chloroperbenzoic acid which gave the *N*-oxide 112. Subsequent treatment of the *N*-oxide 112 with Ac₂O and phosphorus oxychloride according to Example 14 afforded 2-chloro-8-methoxy-quinazoline (113).

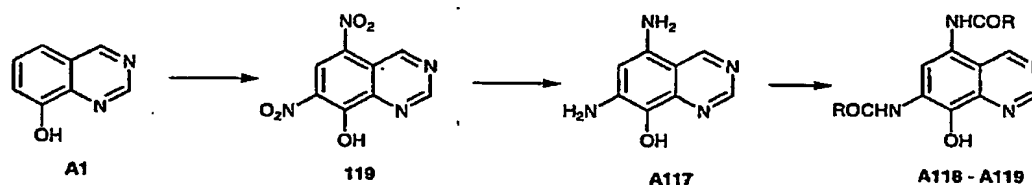
Example 15 Preparation of 2-[(*N'*-methyl)-heterocyclic]-8-hydroxy-quinazolines (A113 – A116) (Scheme A13)

A solution of 2-chloro-8-methoxy-quinazoline (113) (10 mmol) in pyridine (10 mL) was added methylamine hydrochloride (Aldrich, 15 mmol). After 16 h at RT, the mixture was concentrated *in vacuo*. Subsequent column purification of the residue gave the 2-(*N'*-methyl)-compound 114.

Amination of **114** (1 mmol) with 2-bromopyridine (1.2 mmol) in the presence of $[Pd_2(dba)_3]$ and DPPP according to the method described¹¹ in the literature, provided 8-methoxy-2-[(*N'*-pyridyl)methyl]-quinazoline (**115**). Subsequent treatment of **115** with BBr_3 (Example 6) afforded the 8-hydroxy-quinazoline derivative **A113**.

Amination of **114** was repeated using a range of 2-bromo-substituted heterocyclics: 2-bromothiazole, 4-bromo-1*H*-imidazole and 4-bromo-1-methylimidazole which provided **116** - **118**; and subsequent cleavage of the *O*-methyl ether with BBr_3 , gave **A114** - **A116**.

Example 16 Preparation of 5,7-Diamino-8-hydroxy-quinazoline (A117) (Scheme A14)



Scheme A14

To a solution of 8-hydroxy-quinazoline (**A1**) (0.05 mol) in acetic acid (175 mL) was added a solution of nitric acid (0.16 mol) in acetic acid (25 mL), keeping the temperature below 30 °C. After 2 h, the 5,7-dinitro-compound **119** was isolated by filtration, washed with H_2O , and dried.

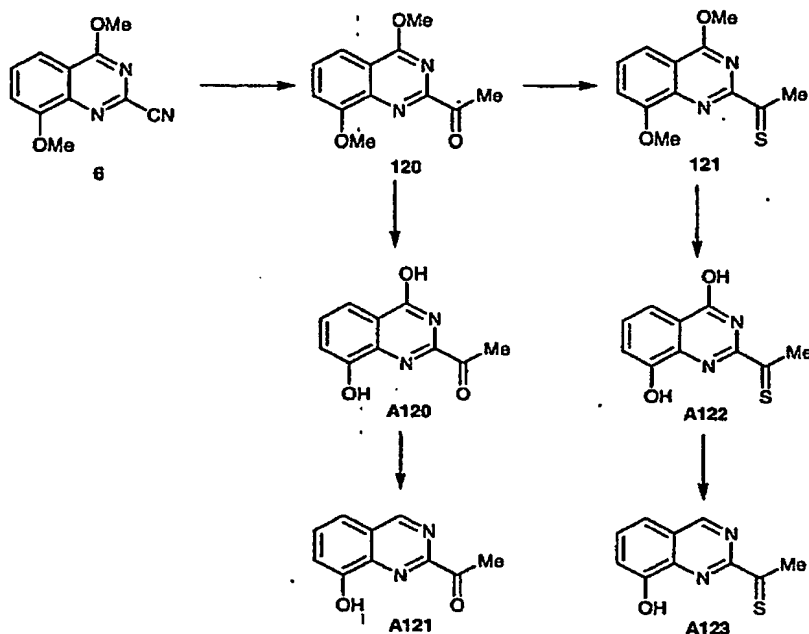
Hydrogenolysis of the 5,7-dinitro-compound **119** (0.045 mol) in MeOH (200 mL) in the presence of platinum oxide gave, after filtration to remove solids and concentration, 5,7-diamino-8-hydroxy-quinazoline (**A117**).

Example 17 Preparation of 5,7-Diacylamino-8-hydroxy-quinazolines (A118 - A119) (Scheme A14)

Acetic acid (2 mmol) and CDI (2.2 mmol) were heated under reflux in dry THF (10 mL) for 1 h. 5,7-Diamino-8-hydroxy-quinazoline (**A117**) (2 mmol) was added and the mixture heated under reflux for 16 h. Removal of volatiles *in vacuo* and subsequent column chromatography of the resulting residue provided 5,7-desacetamido-8-hydroxy-quinazoline (**A118**).

In a similar fashion, treatment of 5,7-diamino-8-hydroxy-quinazoline (**A117**) with benzoic acid provided 5,7-dibenzoylamido-8-hydroxy-quinazoline (**A119**).

Example 18 Preparation of 2-Acetyl-4,8-dihydroxy-quinazoline (A120)
(Scheme A15)



Scheme A15

Methylmagnesium bromide (1.2 mL of a 3 M solution in diethyl ether, 3.5 mmol) was added dropwise into a stirred solution of **6** (0.6 mmol) in diethyl ether (10 mL) at -15°C . The resulting solution was allowed to warm to RT over 2 h and stirred at RT for a further 4 h. The reaction mixture was then quenched with saturated NH_4Cl and extracted with ethyl acetate (10 mL x 3), the extracts combined, dried and concentrated to provide **120**.

Treatment of **120** with BBr_3 (Example 6) gave **A120**.

15 Preparation of 2-Acetyl-8-hydroxy-quinazoline (A121) (Scheme A15)

Successive treatment of **A120** with phosphorus oxychloride, *p*-toluenesulphonhydrazide and Na_2CO_3 (Example 6) gave **A121**.

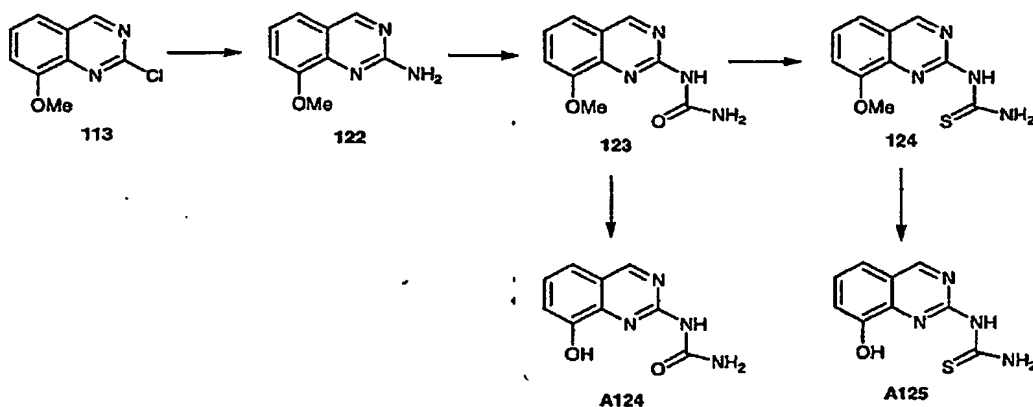
Example 19 Preparation of 2-S-Acetyl-4,8-dihydroxy-quinazoline (A122) (Scheme A15)

A solution of **120** (1 mmol) and Lawesson's reagent (0.7 mmol) in THF (10 mL) was heated under reflux for 16 h and allowed to cool. Concentration and subsequent column chromatography of the residue gave **121**. Treatment of **121** with BBr₃ (Example 6) gave **A122**.

Preparation of 2-S-Acetyl-8-hydroxy-quinazoline (A123) (Scheme A15)

Successive treatment of **A122** with phosphorus oxychloride, *p*-toluenesulphonhydrazide and Na₂CO₃ (Example 6) gave **A123**.

Example 20 Preparation of 8-Hydroxy-quinazoline-2-urea (A124) (Scheme A16)



Scheme A16

The 2-chloride **113** was converted into the amine **122** according to standard Chichibabin reaction conditions. The 2-amine **122** (1 mmol) and isocyanate (1 mmol) in dry CHCl₃ (10 mL) were then heated under reflux for 16 h. This gave, after filtration, the urea derivative **123**. Treatment of **123** with BBr₃ gave **A124**.

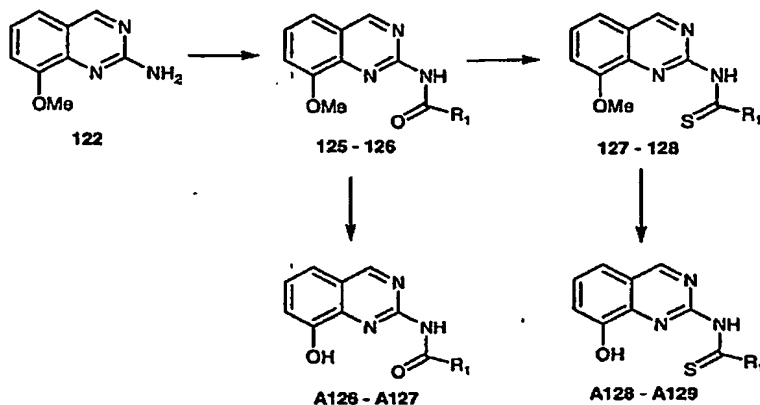
Example 21 Preparation of 8-Hydroxy-quinazoline-2-thiourea (A125) (Scheme A16)

The 2-urea derivative **123** (1 mmol) and Lawesson's reagent (0.7 mmol) in THF (10 mL) was heated under reflux for 16 h, allowed to cool and concentrated.

Subsequent column purification on silica gave **124**. Treatment of **124** with BBr_3 , according to Example 6, then provided **A125**.

Example 22 Preparation of 2-Acylamido-8-hydroxy-quinazolines (A126 – A127)
(Scheme A17)

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Scheme A17

The 2-amine **122** was acylated under standard conditions: Ac_2O gave **125**; benzoic anhydride gave **126**. Respective treatment of **125** and **126** with BBr_3 (Example 6) provided **A126** and **A127**.

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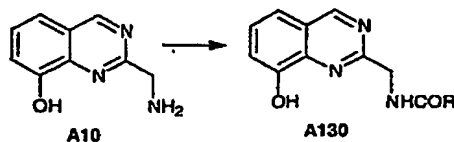
Example 23 Preparation of 2-Thioacylamido-8-hydroxy-quinaxolines
(A128 – A129) (Scheme A17)

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The acylamido compounds **125** and **126** were individually treated with Lawesson's reagent according to conditions previously described (Example 21), which furnished, after standard workup, **127** and **128**. Subsequent *O*-methyl ether cleavage with BBr_3 then gave **A128** and **A129**, respectively.

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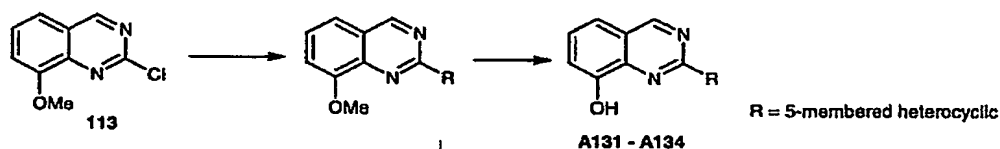
Example 24. Preparation of 2-(acylamido)-methyl-8-hydroxy-quinazolines (A130) (Scheme A18)



Scheme A18

Standard acylation of the amine A10 using a range of acid anhydrides provided, after column purification, the 2-(acylamido)-methyl derivatives A130.

Example 25 Preparation of 2-(Azole)-8-hydroxy-quinazolines (A131 – A134) (Scheme A19)

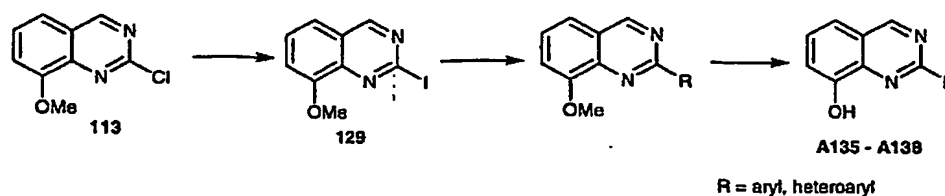


Scheme A19

A mixture of the 2-chloride 113 (0.5 mmol) and pyrazole (2.5 mmol) was heated at 175 °C in a steel autoclave for 48 h. The crude product was then treated with BBr₃ according to the procedure described in Example 6. Subsequent purification by column chromatography gave 2-pyrazol-1-yl-quinazolin-8-ol (A131).

The above procedure was repeated using imidazole, 2-methylimidazole and 1H-1,2,3-triazole to give A132 - A134.

Example 26 Preparation of 2-Aryl(or heterocyclic)-8-hydroxy-quinazolines (A135 – A138) (Scheme A20)



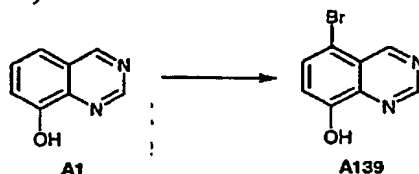
Scheme A20

According to a literature¹² procedure, the 2-chloride **113** (5 mmol) was treated with acetyl chloride and sodium iodide in AcCN. Standard workup followed by column chromatography on silica furnished the 2-iodo-compound **129**.

To a stirred solution of **129** (0.1 mmol) and PdCl₂(PPh₃)₂ (5 mg) in THF (2.5 mL) under an argon atmosphere at RT was added over 5 min 2-pyridylzinc bromide (0.37 mL of a 0.5 M solution in THF, 0.185 mmol). After 2 h, saturated NH₄Cl (5 mL) was added and the mixture extracted with dichloromethane (10 mL x 3). The combined extracts were washed with H₂O and brine, dried and concentrated. Subsequent column chromatography on silica gave 8-methoxy-2-pyrid-2-yl-quinazoline. The 8-*O*-methyl ether was cleaved according to the procedure of Example 6, to give 2-(pyrid-2-yl)-8-hydroxyquinazoline (**A135**).

The reaction was repeated using: 2-(methylthio)phenylzinc iodide, 2-(ethoxycarbonyl)phenylzinc iodide and 6-methylpyridylzinc bromide to give **A136 – A138**.

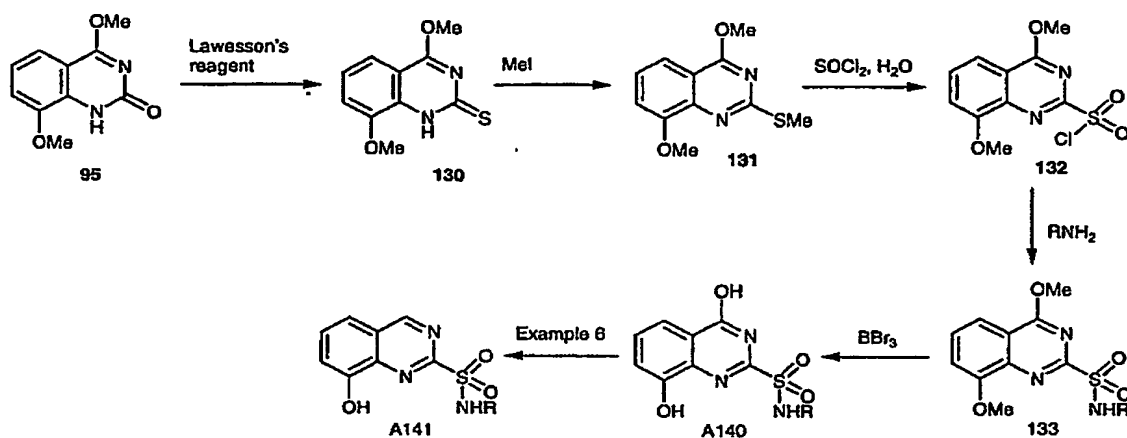
Example 27 Preparation of 5-Bromo-8-hydroxy-quinazoline (A139) (Scheme A21)



Scheme A21

Bromination of 8-hydroxyquinazoline (**A1**) with *N*-bromosuccimide, according to the method previously described by Gerson and McNeil,¹⁵ gave 5-bromo-8-hydroxyquinazoline (**A139**).

**Example 28 Preparation of 8-Hydroxy-quinazoline-2-sulfonic acid
(2-pyridin-2-yl-ethyl)-amide (A140 and A141) (Scheme A22)**

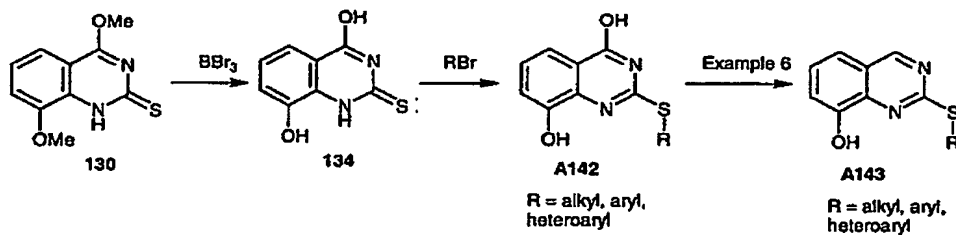


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Scheme A22

RNH₂: preferably, ethylamine, histamine, 2-(2-aminoethyl)pyridine, 2-(2-methylaminoethyl)pyridine

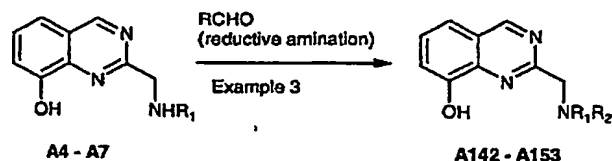
10 Example 29 Preparation of 2-Alkyl(or aryl or heterocyclic)sulfanyl-8-hydroxy-quinazoline (A142 and A143) (Scheme A23)



Scheme A23

15 RBr: preferably, 2-bromopyridine, 2-bromothiazole, 4-bromo-1H-imidazole, 4-bromo-1-methylimidazole

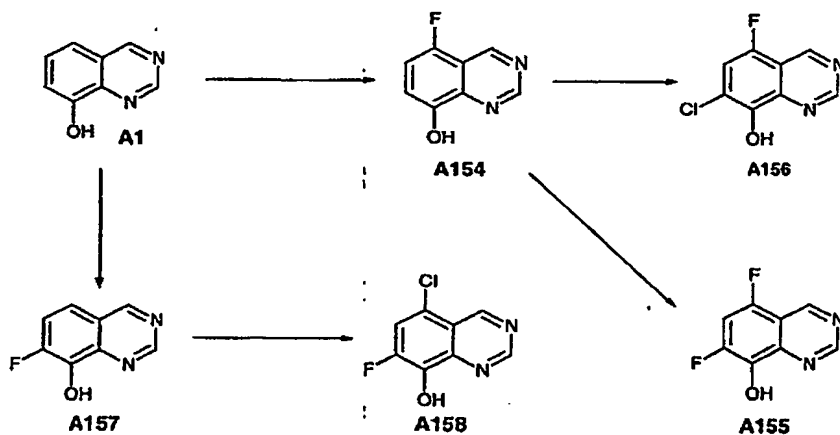
Example 30 Reductive amination of amines (A4 – A7) from Example 3 to give A142 – A153 (Scheme A24)



Scheme A24

RCHO: preferably, 4-imidazolecarboxaldehyde, 2-thiazolecarboxaldehyde, 2-pyridinecarboxaldehyde

Example 31 Preparation of 5-fluoro-, 7-fluoro, 5,7-difluoro, 5-chloro-7-fluoro and 7-chloro-5-fluoro-quinazolines (A154 – A158) (Scheme A25)



Scheme A25

The synthesis of the fluorinated quinazoline derivatives (A154 – A158) followed literature methods¹⁸ for 8-hydroxy-quinoline.

Example 32 Preparation of 8-Hydroxy-quinazoline-7-carboxylic acid amides (A160) (Scheme A26)



Scheme A26

8-Hydroxy-quinazoline (**A1**) was converted into 8-hydroxy-quinazoline-2-carboxylic acid (**A159**) following the literature method²⁰ for the carboxylation of 8-hydroxy-quinoline. Subsequent conversion of **A159** into the amide **A160** followed the procedure described in Example 8.

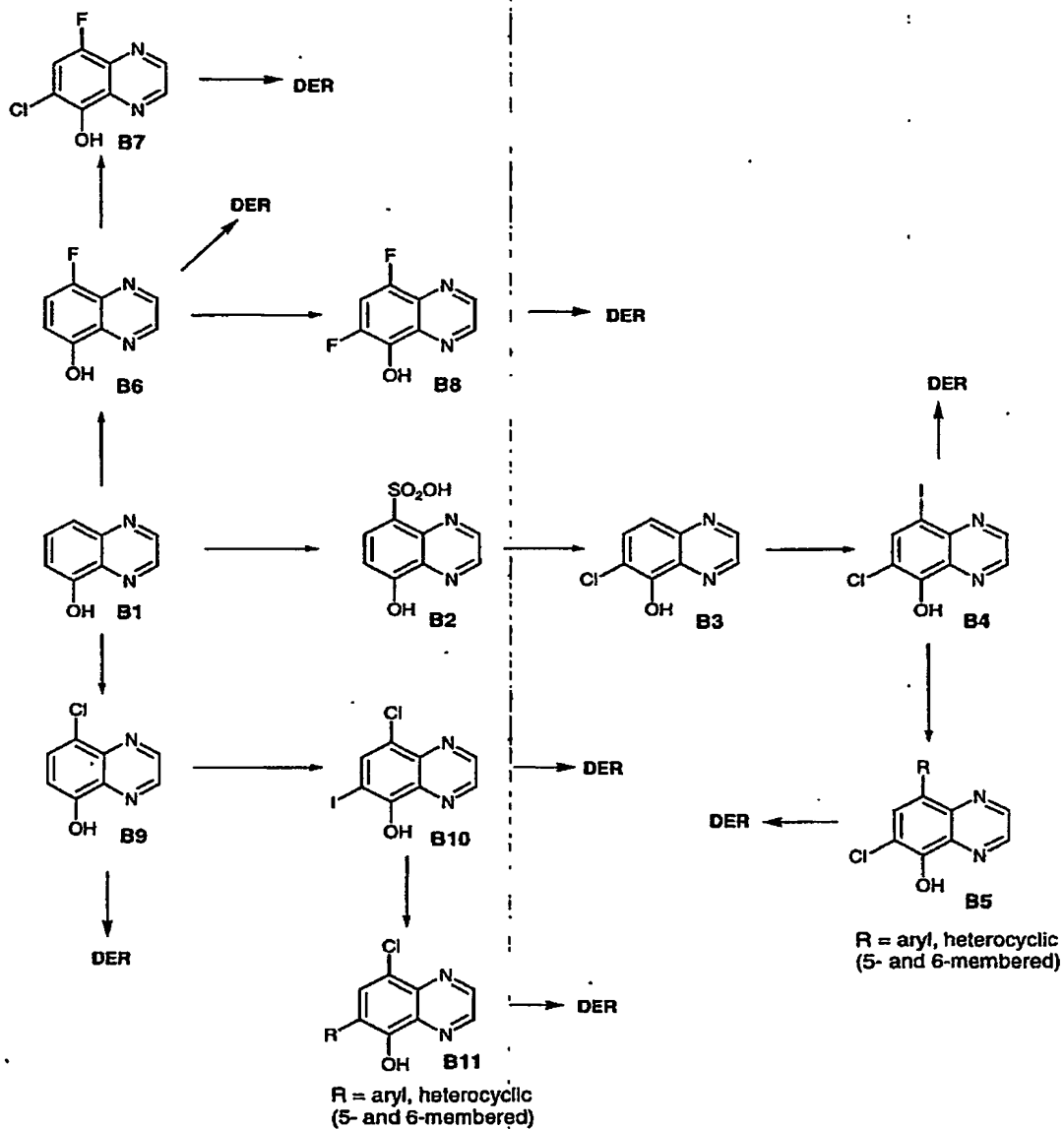
5

PART B: SYNTHESIS OF 8-HYDROXY-QUINOXALINE DERIVATIVES

Charts B1 – B4 show the routes to the 8-hydroxy-quinoxaline derivatives. The synthetic procedures were, unless otherwise indicated, analogous to those previously described in

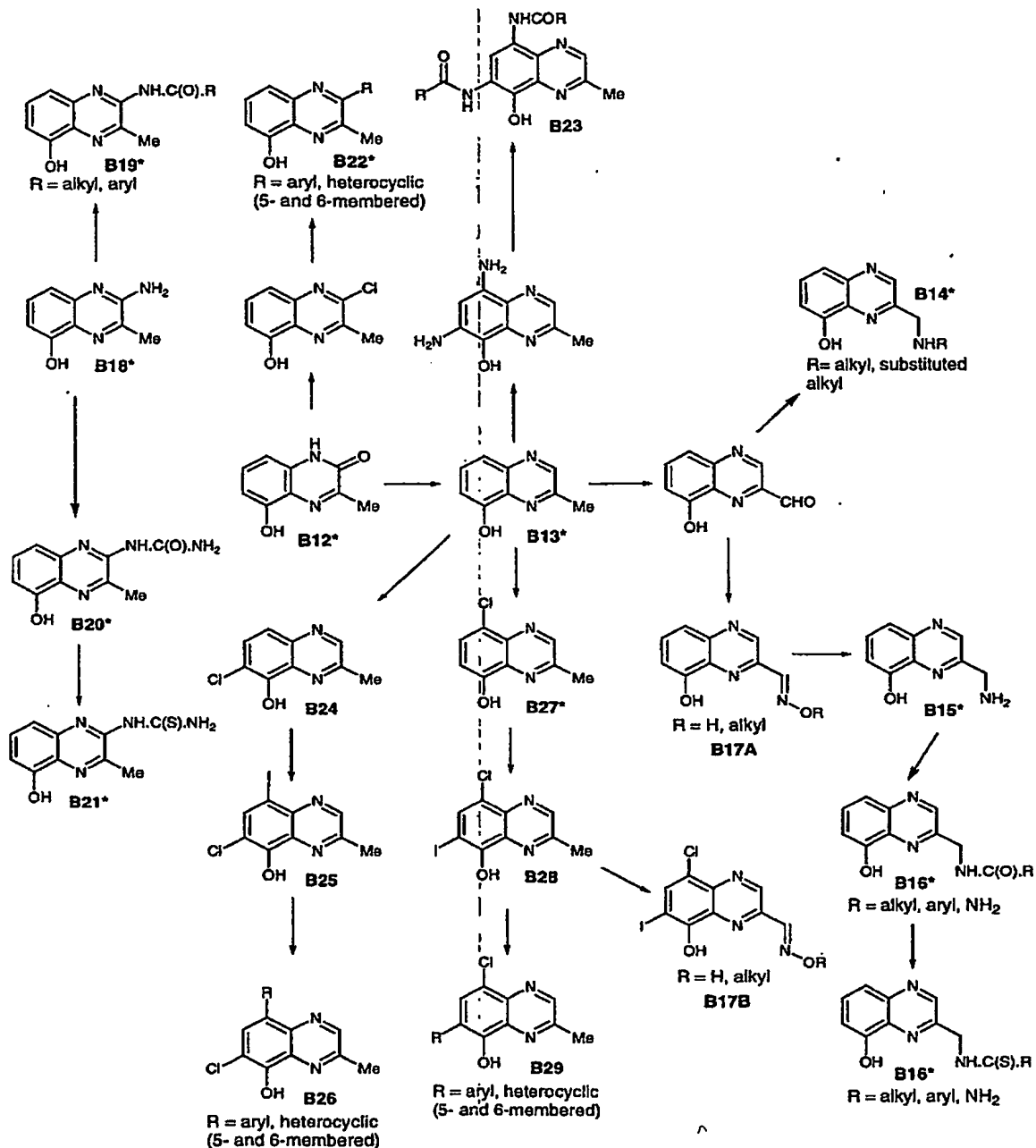
10 Part A.

CHART B1



DER: derivatives via substitution of 2- and /or 3-chloro/cyano group(s) to a subject in need thereof e.g. NR_1R_2 ($\text{R}_1 = \text{H}$, alkyl; $\text{R}_2 = \text{alkyl, aryl, heterocyclic}$), SR , CONHR , alkylamino

- 51 -
CHART B2



DER: derivatives via substitution of 3-chloro/cyano group to a subject in need thereof e.g. NR₁R₂ (R₁ = H, alkyl; R₂ = alkyl, aryl, heterocyclic), SR, CONHR, alkylamino

* - and halogen-substitution, preferably at the 5- and/or 7-position(s)

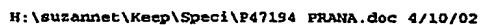
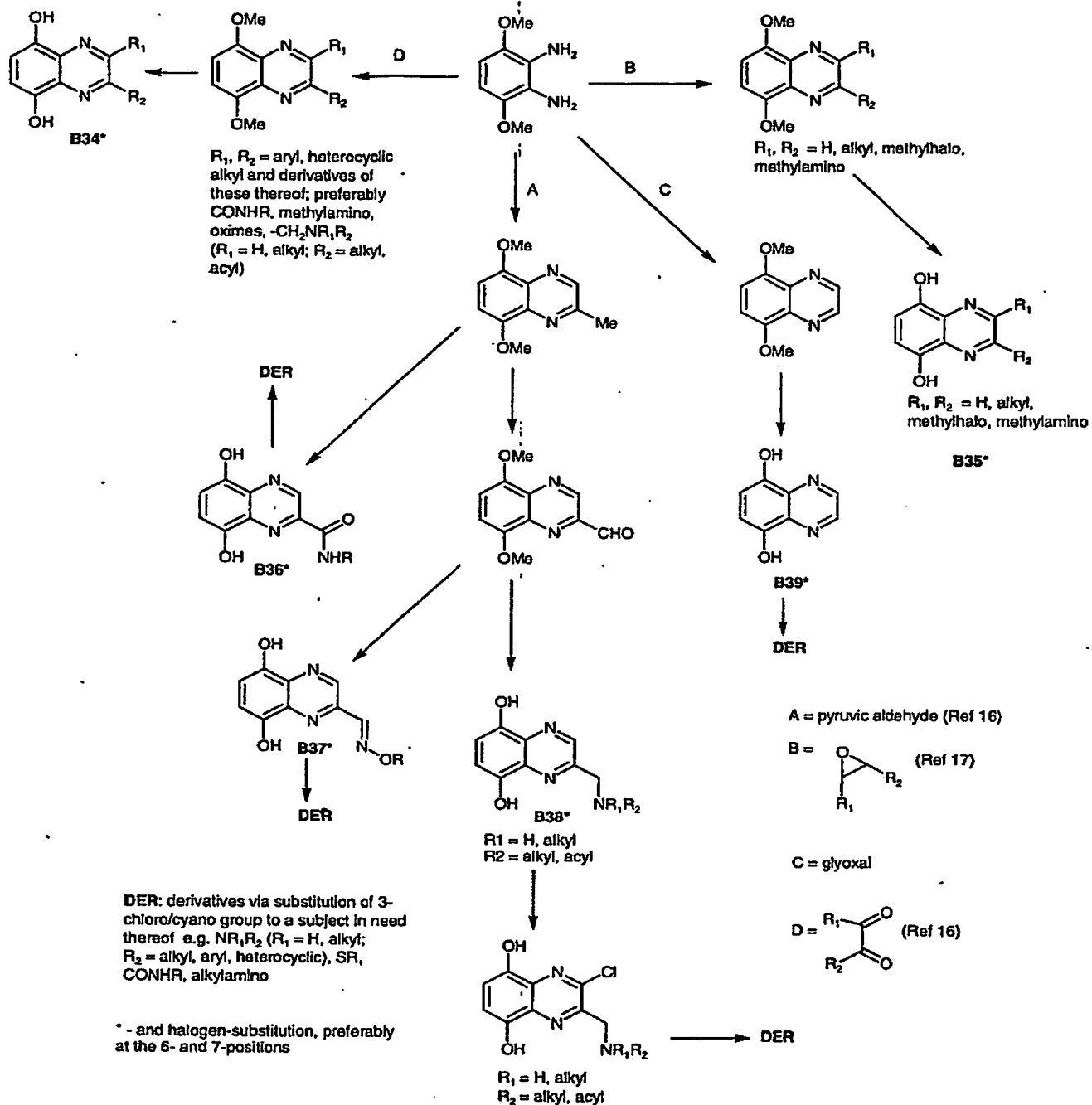


CHART B4



PART C: SYNTHESIS OF 8-HYDROXY-CINNOLINE DERIVATIVES

Charts C1 and C2 show the routes to the 8-hydroxy-cinnoline derivatives. The synthetic procedures were, unless otherwise stated, analogous to those previously described in Part A.

CHART C1

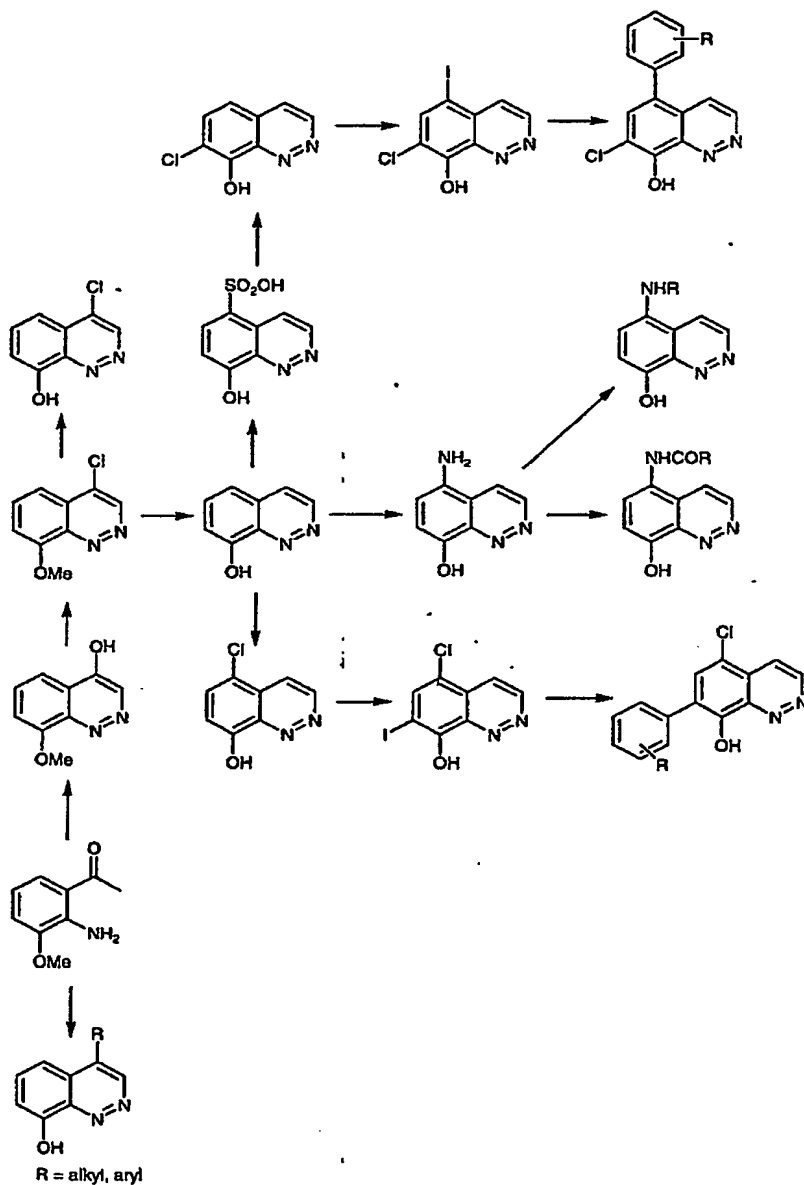
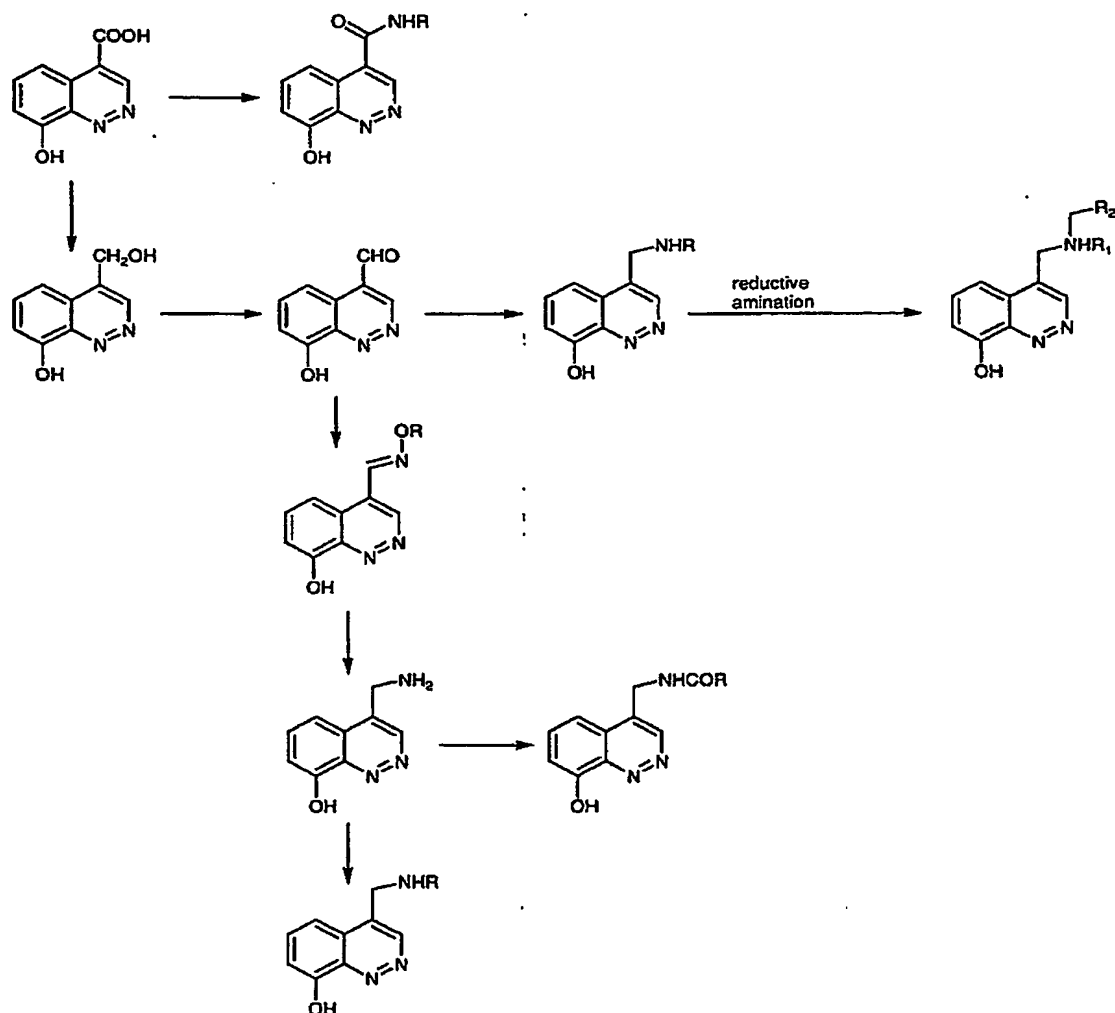


CHART C2



PART D: SYNTHESIS OF 4,7(4,10)-PHENANTHROLIN-5-OL DERIVATIVES

5

Charts D1 – D4 show the routes to the 4,7(4,10)-phenanthroline-5-ol derivatives. The synthetic procedures were, unless otherwise stated, analogous to those previously described in Part A. The reactions shown in Chart D1 were also repeated using 4,7-phenanthroline-5,6-diol (**D3**) instead of 4,7-phenanthroline-5-ol (**D1**) as starting material.

CHART D1

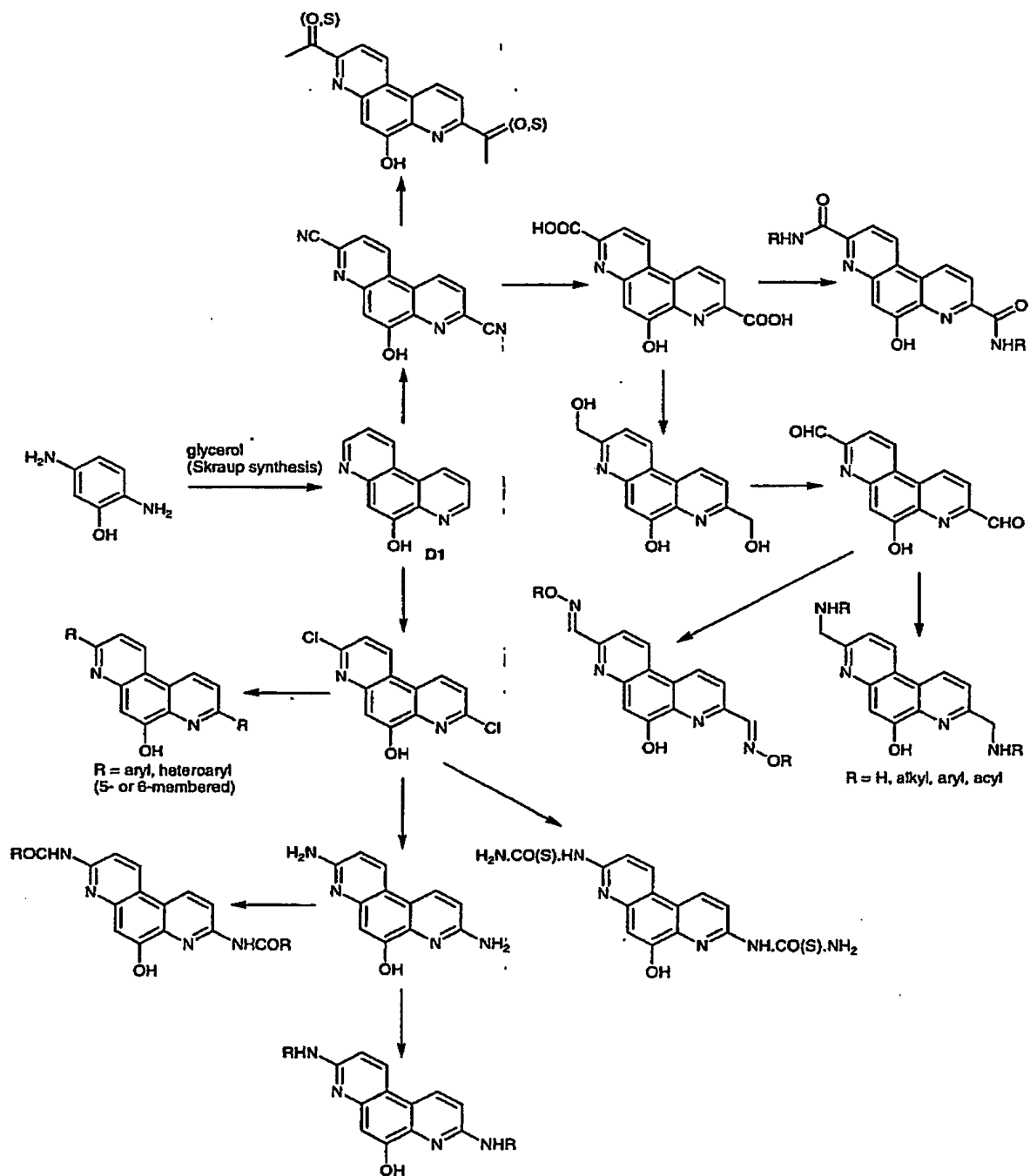


CHART D2

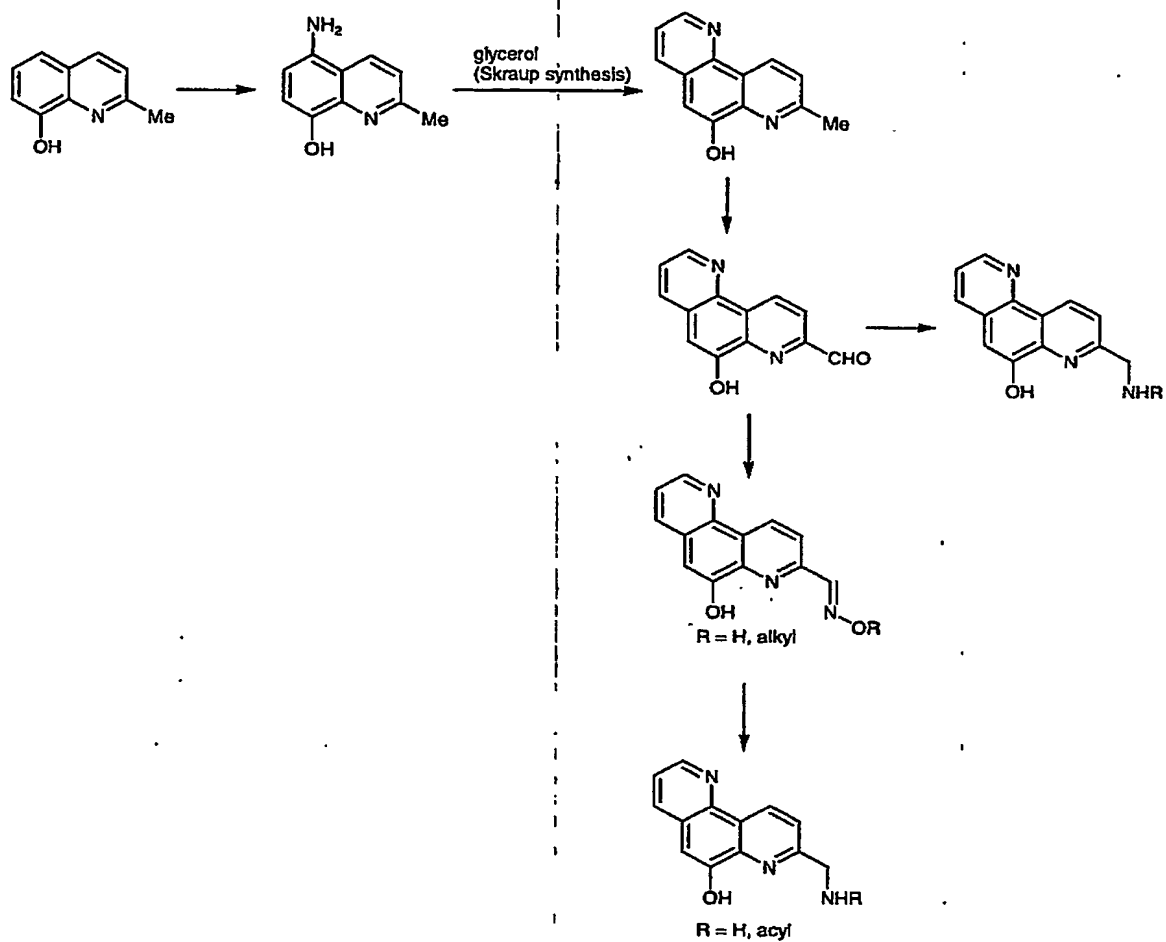
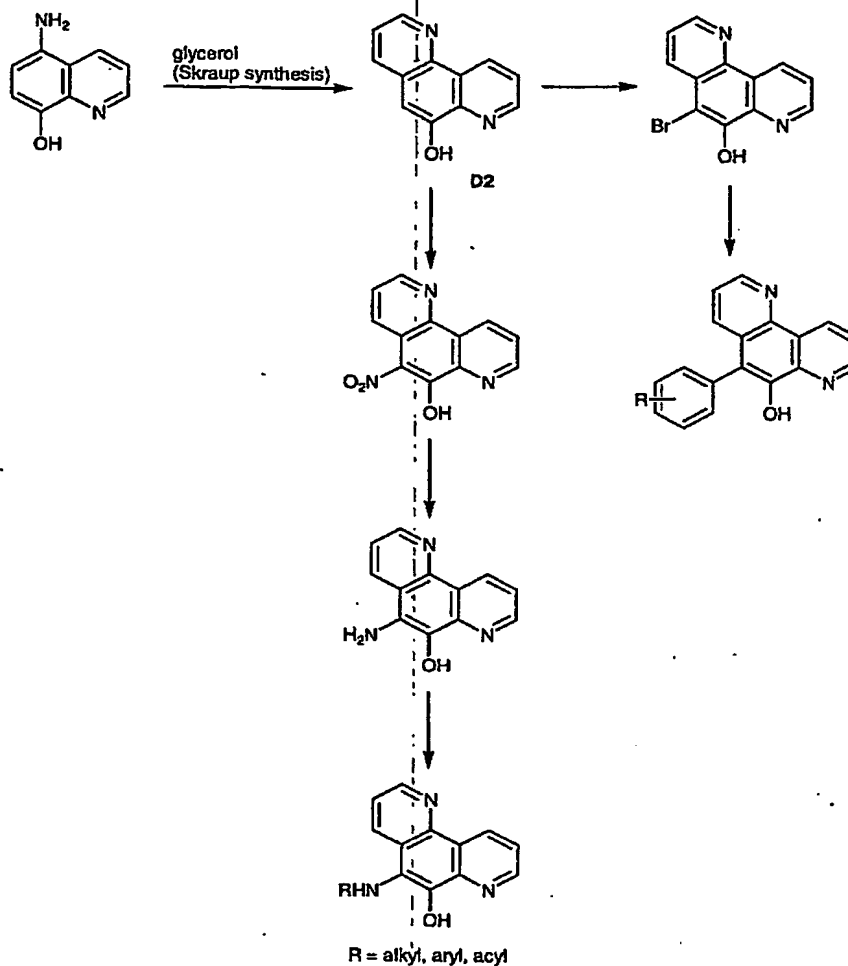


CHART D4

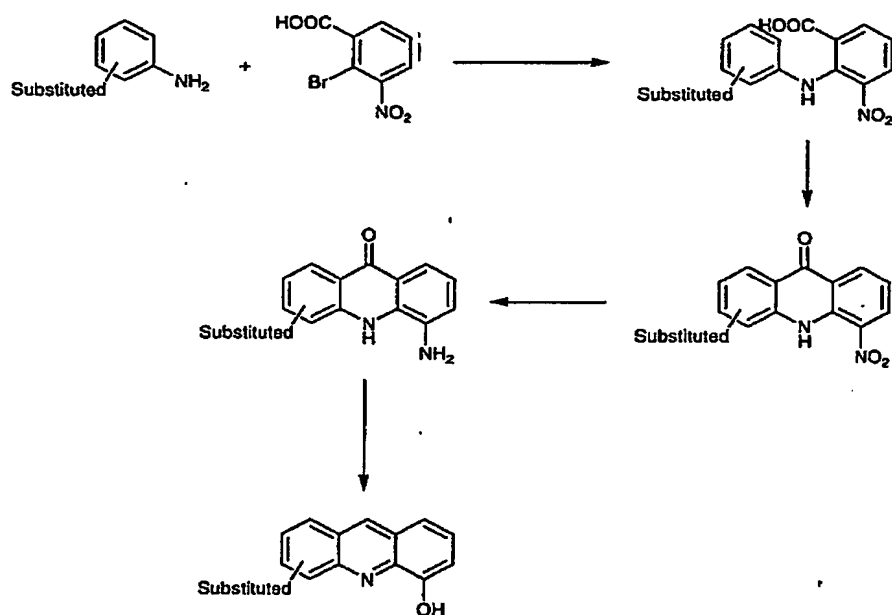
**Example Skraup synthesis of 4,7-Phenanthroline-5-ol (D1)**

- 5 A stirring mixture of 3-hydroxy-*p*-phenylenediamine (0.185 mol),
 glycerol (1.17 mol), arsenic solution (100 mL; prepared from 123 g of arsenic
 pentoxide in 100 mL H₂O) and diluted sulphuric acid (400 mL; prepared by adding 240
 mL of concentrated sulphuric acid to 200 mL H₂O) was heated under reflux for 4 h,
 allowed to cool and then made alkaline with concentrated ammonia. The mixture was
 10 extracted with benzene. Removal of the benzene afforded 4,7-Phenanthroline-5-ol (D1).

PART E: SYNTHESIS OF 4-HYDROXY-ACRIDINE DERIVATIVES

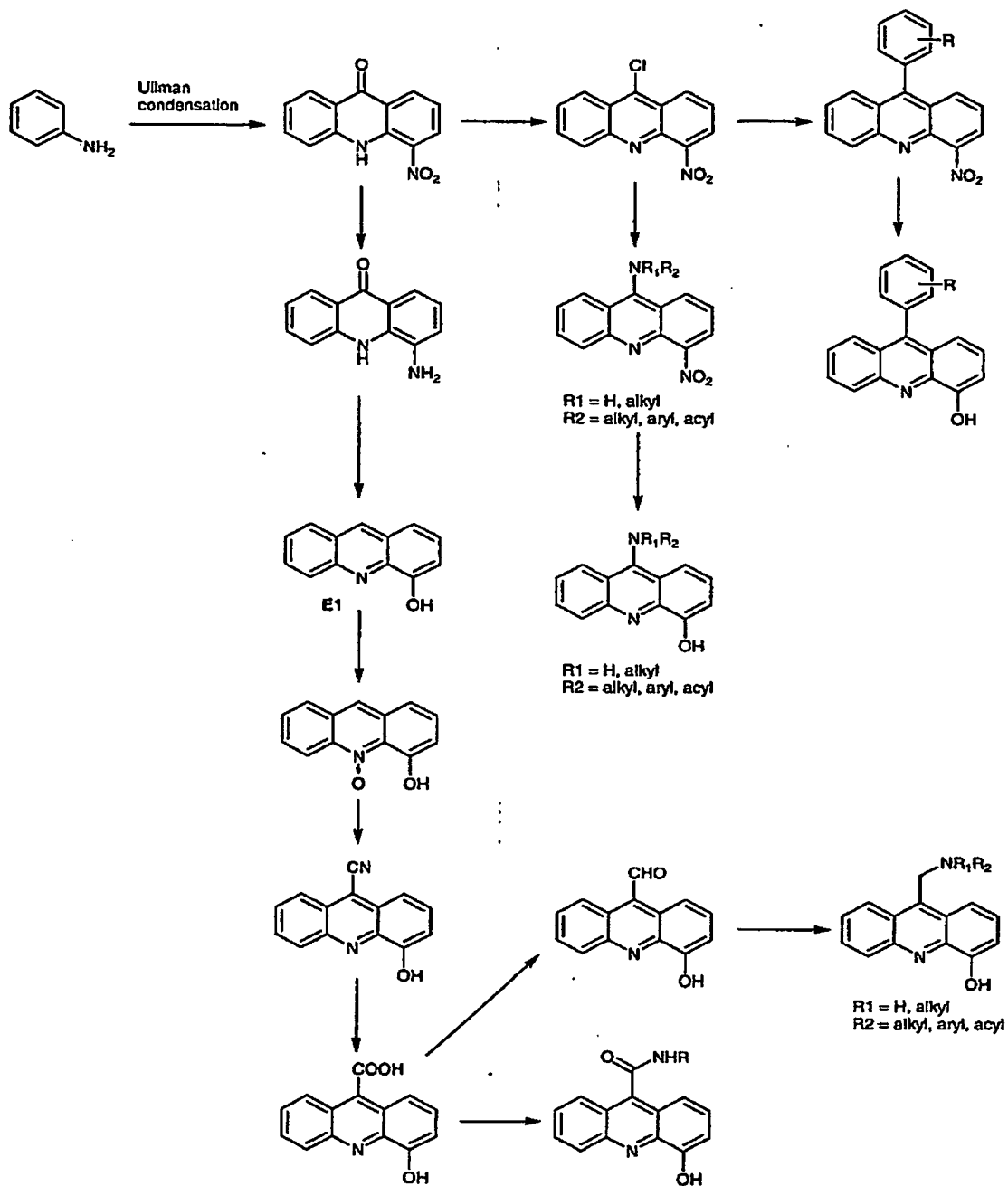
The 4-hydroxy-acridines were prepared via Ullman condensation^{4,14} of a substituted 2-halobenzoic acid and a substituted aniline as shown in Chart E1. Hence, condensation of aniline itself with 2-bromo-3-nitro-benzoic acid gave 4-hydroxy-acridine (E1). In an analogous fashion, *o*-anisidine gave 4-amino-5-hydroxy-acridine and 4,5-dihydroxy-acridine. Further derivatives of these acridines (Chart E2) were prepared using analogous reaction conditions previously described for the synthesis of compounds in Parts A – D.

CHART E1



Substituents on aniline include alkyl, methoxy, halogen

CHART E2



The compounds, 4,5- and 4,8-dihydroxy-phenazines (**F1** and **F2**), were prepared according to the literature³ procedure. The synthesis of derivatives of these 5 compounds, as summarized in Charts F1 – F3, unless otherwise stated, followed reaction conditions analogous to those previously described in Parts A – E.

CHART F1

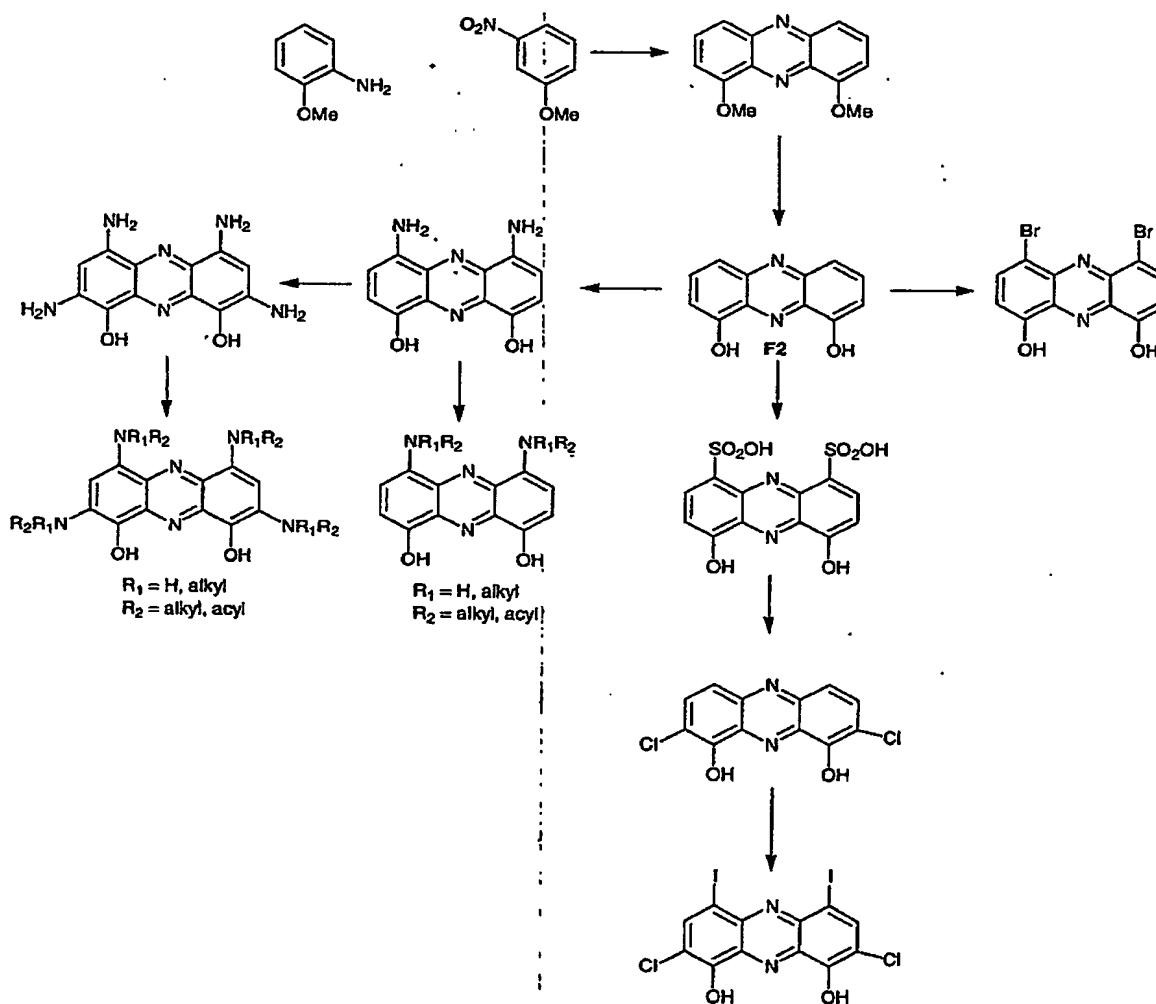
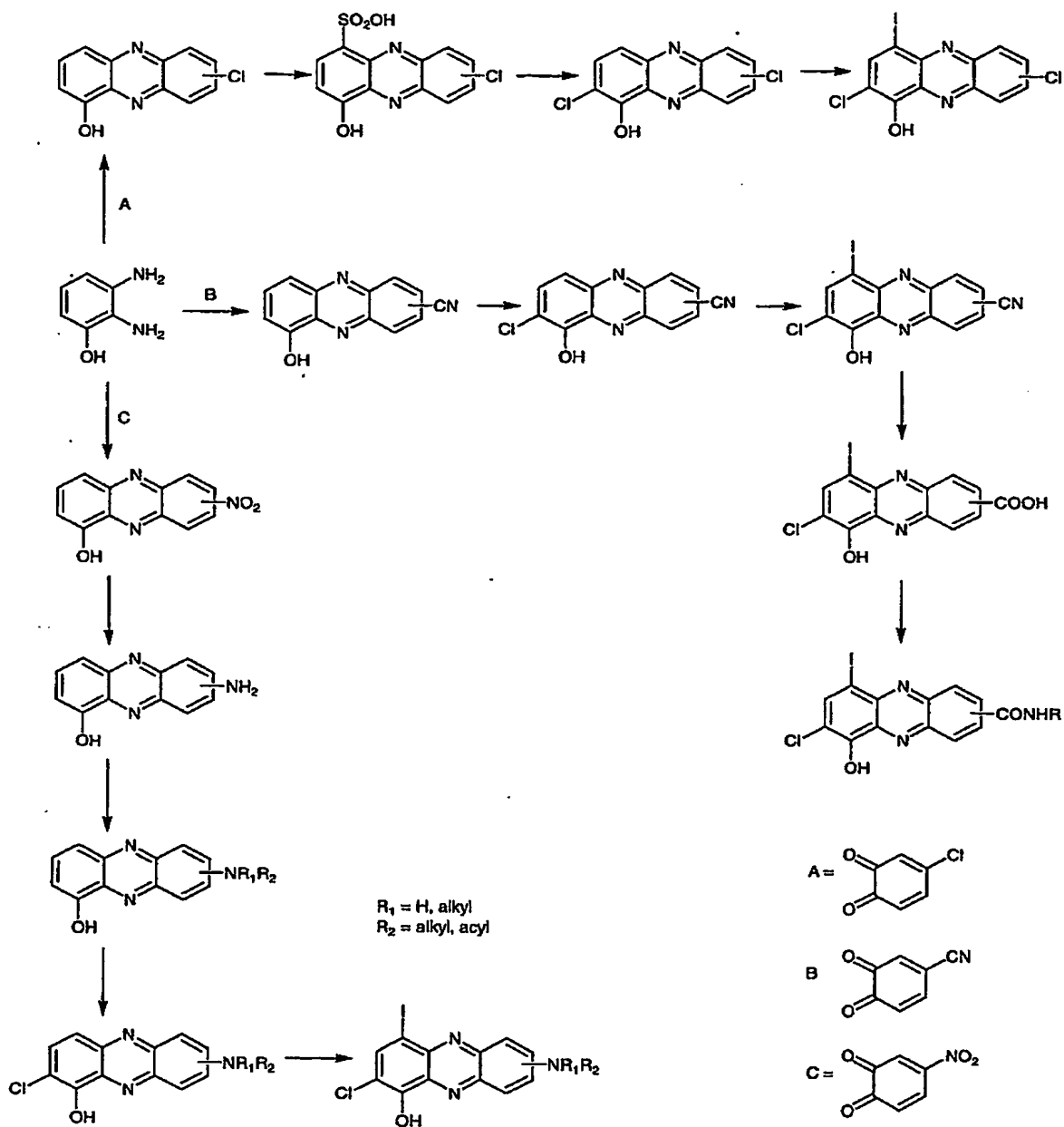




CHART F3



5 The following assays are used in the assessment of new compounds according to the invention for suitability for use in the methods of the invention.

Fluorometric H₂O₂ assay

A fluorometric assay is used to test for the ability of a test compound to inhibit hydrogen peroxide generation by A β in the presence of copper based on dichlorofluorescein diacetate (DCF; Molecular Probes, Eugene OR). The DCF solution (5mM) in 100% dimethyl sulphoxide (previously purged with argon for 2hr at 20°C) is deacetylated in the presence of 0.25M NaOH for 30min and neutralised at pH 7.4 to a final concentration of 1mM. Horseradish peroxidase(HRP) stock solution is prepared to 1 μ M at pH 7.4. The reactions are carried out in PBS, pH 7.4 in a 96 well plate (total volume =250 μ l/well). The reactions solutions contain A β 1-42 at concentrations which may be in the range 50nM to 1 μ M, copper-glycine chelate (Cu-Gly, prepared by adding CuCl₂ to glycine in the ratio of 1:6 and added to the A β in the proportion 2Cu-Gly : 1A β), reducing agents including dopamine (5 μ M) or ascorbic acid, deacetylated DCF 100 μ M, and HRP, 0.1 μ M. 1-10 μ M EDTA or another chelator may also be present as a control for free copper, but is not required for the assay to function. The reaction mixture is incubated at 37C for 60 min. Catalase (4000 units/ml) and H₂O₂ (1-2.5 μ M) standards in PBS pH 7.4 may be included as positive controls. Fluorescence is recorded using a plate reader with excitation and emission filters at 485nM and 530nM respectively. H₂O₂ concentration may be established by comparing fluorescence with the H₂O₂ standards. Inhibition of A β H₂O₂ production is assayed by including a given concentration of test compound(s) in the test wells.

Neurotoxicity Assays

Primary cortical neuronal cultures

Cortical cultures are prepared as previously described (White *et al.*, 1998). Embryonic day 14 BL6Jx129sv mouse cortices are removed, dissected free of meninges and dissociated in 0.025% (wt/vol) trypsin. Dissociated cells are plated in 24 well culture plates (Greiner GmbH, Austria) at a density of 2 x 10⁶ cells/mL in MEM with 10% (vol/vol) FCS and 10% (vol/vol) HS. Cultures are maintained at 37°C in 5% CO₂. Prior to experiments, the culture medium is replaced with MEM plus N2 supplements.

Primary cerebellar granule neuronal cultures

Cerebella from post-natal day 5-6 (P5-6) mice are removed and dissected free of meninges and dissociated in 0.025% trypsin. Cerebellar granule neurons (CGN) are plated in 24 well culture plates at 350 000 cells/cm² in BME (Gibco BRL) supplemented with 10% FCS, 2 mM glutamine and 25 mM KCl. Gentamycin sulphate

(100 µg/mL) is added to all plating media and cultures are maintained at 37°C in 5% CO₂.

Assays for cell viability

5 MTT assay for cell viability

Cell viability is determined using the MTT assay. Culture medium is replaced with 0.6 mg/mL MTT in control salt solution (Locke's buffer containing 154 mM NaCl, 5.6 mM KCl, 2.3 mM CaCl₂, 1.0 mM MgCl₂, 3.6 mM NaHCO₃, 5 mM HEPES and 5.6 mM glucose, pH 7.4) for 30 min. The MTT is removed and cells
10 solubilized with dimethyl sulfoxide. 100 µL aliquots are measured with a spectrophotometer at 570 nm.

LDH assay for cell viability

Cell death is determined from culture supernatants free of serum and cell
15 debris using the lactate dehydrogenase (LDH) Cytotoxicity Detection Kit (Boehringer Ingelheim) according to the manufacturer's instructions.

Assay for neurotoxicity at low Aβ concentration

Cortical cells are prepared following the protocol of White et al. (1998),
20 with the following modifications:

A) On the 5-6th day the medium is changed for Neurobasal medium plus B27 but minus antioxidant.

B) On the 8-9 day the medium is replaced by medium
containing test reagents, including Aβ (200-1000 nM),
25 Cu-Gly (400-2000 nM) and dopamine (5-20 µM in PBS).
EDTA (10 µM in PBS) is included throughout to eliminate undesired reactions between free copper and dopamine. However, when testing new drugs, it is advisable not to include EDTA in the Aβ-Cu-Dopamine mixture. For controls, the dopamine volume is replaced with PBS 7.4; the Cu-Gly volume is replaced with water and the Aβ volume is
30 replaced with water.

Aβ peptide solution is prepared by dissolving the peptide in water and centrifuge at 13,000 rpm, for 3-5 min. The supernatant is carefully harvested and its concentration measured by absorbance at 214 nm using the absorbance standard curve. The following is the mixture sequence and example of approx. volumes of each
35 compound:

For a final volume of 1000 µL, the following sequence is carried out:

A β is added using 6.3 μ L of A β stock (80 μ M), to give a final concentration of 500 nM. Thereafter 10 μ L of Cu-Gly stock (100 μ M) is added to give a final concentration of 1000 nM. 68.7 μ L of H₂O and 10 μ L of EDTA 1 mM are added, to give a final concentration 10 μ M of EDTA. 900 μ L of Neurobasal medium plus B27 without
5 antioxidant or Locke's buffer is then added and the solution is mixed. 5 μ L of freshly made Dopamine stock (1 mM) is then added to give a final dopamine concentration 5 μ M, and the solution is mixed again. The cell medium in each well of the culture is replaced with 250 μ L of the mixture, and the cultures are incubated for 16-24 h (37° C).
10 Following incubation, each well is gently washed twice with Locke's buffer, and then the Locke's buffer is replaced with Neurobasal medium (250 μ L). Three empty wells are included as background controls.
25 μ L of MTS stock is added to each well and incubated for 2-4 hrs at 37° C. The absorbance is then read at 490nm.

15 *Caspase assay*

To measure caspase activity in neuronal cultures, growth medium is removed, cells are washed twice with control salt solution (pH 7.4) and ice-cold cell extraction buffer is added directly to the cultures. The extraction buffer consists of 20
20 mM Tris (pH 7.4), 1 mM sucrose, 0.25 mM EDTA, 1 mM dithiothreitol (DTT), 0.5 mM PMSF, 1% Triton X-100 (Tx-100) and 1 μ g/mL of pepstatin and aprotinin. After incubation for 15 min on ice, the extraction buffer is removed, centrifuged for 5 min at 4°C in a microcentrifuge and 100 μ L of supernatant is added to each well of a 96 well plate. 100 μ L of 200 μ M substrate (either DEVD-pNA, VEID-pNA or IETD-pNA for caspases 3, 6 and 8 respectively) is added to each well to give a final concentration of
25 100 μ M substrate. Plates are incubated at 37°C for 2, 4, 6 or 24 hr and the absorbance is determined at a wavelength of 415 nm (Abs415). The absorbance reading is compared to a known standard of pNA alone.

Annexin V assay

30 To determine the level of annexin V binding to cells, cultures are washed twice with control salt solution (pH 7.4) followed by the addition of annexin V-FITC at a concentration of approximately 0.5 μ g/mL in control salt solution (pH 7.4). Propidium iodide (10 μ g/mL) is also added to the cultures at the same time. Cells are incubated in
the dark for 30 min at ambient temperature and subsequently washed three times with
35 fresh control salt solution. Analysis of FITC fluorescence (ex. 488 nm, em. 510 nm) is determined using a Leica DMIRB microscope. Photographs are taken with a Leica MPS

60 camera attachment using ASA400 colour film, and negatives are scanned into Adobe Photoshop v2.0.1.

Lipoprotein oxidation assay

5 Two different assays of metal-mediated lipid peroxidation can be utilized. The first assay involves measuring the oxidative activity of metallated proteins. This is determined by mixing dialyzed metallated or native protein (at designated concentrations) with 0.5 mg/mL LDL for 24 hr (37°C). Lipid peroxidation (LPO) is measured using a lipid peroxidation assay kit (LPO 486, Oxis International Inc.
10 Portland, OR) as per kit instructions. The level of LPO is determined by comparing absorbance (486 nm) with LDL alone (100% LPO). The second assay is used to measure the LPO activity of native proteins in the presence of free, non-protein-bound Cu. This involves adding non-metallated peptides (140 µM) to 0.5 mg/mL LDL together with 20 µM Cu-gly and assaying for LPO as for the metallated proteins. The
15 level of LPO is determined by comparing the absorbance (486 nm) with LDL + Cu-gly (100% LPO). As a negative control, LDL is also exposed to dialysed Cu-gly solutions comparable to those used to Cu-metallate the proteins.

Cytotoxicity induced by Cu-metallated proteins

20 Proteins or synthetic peptides are mixed with metal-glycine solutions at equimolar or two-fold metal to protein concentration. Metal-protein mixtures are incubated overnight at 37°C and then extensively dialysed (24 hr against two changes of dH₂O (3 L/change) at room temperature) using mini-dialysis cups with a 3,500 kilodalton cut-off (Pierce, Rockford, IL). Dialysis of proteins against PBS pH 7.4
25 resulted in metallated proteins with identical activity to dH₂O dialysis.

To determine their neurotoxic effects, metallated proteins, native proteins or peptides are added to two day-old primary cortical neuronal cultures. The cultures are also exposed to Cu-gly (5 or 10 µM) or LDL. Positive control cultures are treated with Cu-gly + LDL or the LPO product, 4-hydroxy-nonenol (HNE, Sigma
30 Chemicals). Cultures are assayed for cell death using the lactate dehydrogenase (LDH) assay kit (Roche Molecular Biochemicals, Nunawading, Australia) according to the manufacturer's instructions.

Acridine orange assay for Aβ-mediated loss of lysosomal acidification

35 Cultured mouse cortical neurons are treated with Aβ1-42 (20µM) for 16 h and then stained with 5 mg/ml acridine orange (AO) for 5 min at 37°C. 15 min at 37°C. The AO-induced fluorescence is measured with a red filter on a fluorescence

microscope. AO is a lysosomotropic weak base which accumulates in the endosomal/lysosomal compartments and displays orange fluorescence during incubation. AO is sequestered inside the lysosomes as long as there is a substantial proton gradient over the lysosomal membranes. Treatment of cells with A β 1-42
5 disrupts the lysosomal membrane proton gradient and relocalises AO into the cytosol, as indicated by the loss of orange fluorescence within 16-24 hr.

Brain amyloid solubilisation assay

This assay is performed in order to assess the ability of a test compound
10 to mobilise A β from the insoluble to the soluble phase of an extract of tissue from *post mortem* human AD brain.

Up to 0.5 g of plaque-bearing cortex without meninges is homogenized using a DIAX 900 homogenizer (Heudolph and Co, Kelheim, Germany) or other
15 suitable device for three 30-second periods at full speed in 2 ml of ice-cold phosphate-buffered saline, pH 7.4. To obtain the phosphate-buffered saline-extractable fraction, the homogenate is centrifuged at 100,000 x g for 30 min and the supernatant removed. Supernatant, either freeze-dried and resuspended or in unconcentrated form, is
20 dissolved in 200 μ l of Tris-Tricine sodium dodecyl sulfate (SDS) sample buffer pH 8.3 containing 8% SDS, 10% 2-mercaptoethanol. Aliquots (10 μ l) are then boiled for 10 minutes before SDS-polyacrylamide gel electrophoresis. The insoluble fraction of the
cortical samples is obtained by resuspending the initial pelleted sample in 1 ml of phosphate-buffered saline. A 50- μ l aliquot of this suspension is then boiled in 200 μ l of sample buffer as above.

Tris-Tricine polyacrylamide gel electrophoresis is performed by loading
25 appropriately diluted samples on to 10% to 20% gradient gels (Novex, San Diego, CA) followed by transfer on to 0.2- μ m nitrocellulose membrane (Bio-Rad, Hercules, CA). A β is detected by using monoclonal antibody W02, which detects residues 5 through 8, 17 (or another suitable antibody) in conjunction with horseradish peroxidase-
30 conjugated rabbit anti-mouse IgG (Dako, Denmark), and visualized by using enhanced chemiluminescence (eg ECL; Amersham Life Science, Buckinghamshire, UK). Each gel includes three lanes containing 0.5, 1, and 2 ng of synthetic A β ₄₀ (Keck
Laboratory, Yale University, New Haven, CT) as reference standards.

Blot films are scanned by using a suitable imaging system such as the
UVP gel documentation system, and densitometry performed using suitable software,
35 eg UVP Labworks . The dynamic range of the film/scanner is determined by using a step tablet (No. 911ST600, Kodak, Rochester NY), a calibrated film exposed by the manufacturer to provide steps of known increasing intensity. The quantifiable range of

signal intensity for densitometric analysis of the mono- and dimeric A β bands is based on the comparison with a curve obtained by scanning and densitometry of the step tablet. Samples in which the signal intensity is low after preliminary assay may be re-assayed by using synthetic standards of lower or higher concentration.

5 All samples are analyzed at least twice, and gel loadings and dilutions are adjusted to fit within the quantifiable region of the standard curve. The proportion of soluble to insoluble A β may be used to determine the efficiency of extraction of the test compound compared with the efficiency of a known compound, such as bathocuproine or clioquinol.

10 *Metal partitioning*

To assay effects upon the partitioning of various metals, including zinc and copper, following extraction of brain tissue in the presence of a test compound, soluble and insoluble fractions from an extract of human brain tissue are prepared as for
15 the amyloid solubilisation assay. Metals in the two fractions are analysed by inductively-coupled plasma mass spectrometry, following appropriate pretreatment with nitric acid and/or hydrogen peroxide where necessary.

Effect of administration of agents on A β deposits in transgenic animals

20 Transgenic mouse models are available for a number of neurological disorders, including Alzheimer's disease (Games et al., 1995; Hsiao et al., 1996); Parkinson's disease (Masliah et al., 2000); familial amyotrophic lateral sclerosis (ALS) (Gurney et al., 1994); Huntington's disease (Reddy et al., 1998); and Creutzfeld-Jakob disease (CJD) (Telling et al., 1994). We have found that one of the transgenic models for
25 Alzheimer's disease, the APP2576 tg mouse (Hsiao et al., 1996) also has a high incidence of cataract. These animal models are suitable for testing the methods of the invention.

Transgenic mice of the strain APP2576 (Hsiao et al 1996) are used. Eight to nine month old female mice are selected and divided into groups for treatment.

30 Mice are sacrificed at intervals, and their brains examined to determine whether the treatment decreases brain amyloid formation, and to identify the most effective administration protocol. The levels of soluble and insoluble A β in the brain and serum are determined using calibrated Western blots. The A β plaque burden in the brain is examined immunohistochemically.

35 Other mice in each group are tested over a period of up to eight months for cognitive performance, using a Morris water maze according to standard methods. The general health and well-being of the animals is also measured every day by a blinded

operator, using a five point integer scale which subjectively rates a combination of features, including motor activity, alertness and general health signs.

Solubility Assay

5 Stock solutions of compounds of formula I or II (1mM) were prepared in dimethyl sulfoxide. Compounds which did not dissolve were classed as not soluble (N). The DMSO stock solutions were diluted 1 in 100 into PBS pH 7.4. Compounds which gave a clear solution were classed as soluble (Y), while those compounds which gave a translucent suspension after dissolution in DMSO were classed as "crashed out" (C).

10

Example 13 - Screening Tests of compound of formula I or II for the treatment of Alzheimer's disease

15

Example 14 - Clinical trial of compound of formula I or II for the treatment of Alzheimer's disease

20 A Phase II clinical trial of the compound of formula I or II for the treatment of AD has been prepared. Inclusion criteria for this Phase II study targeted a region on the ADAS-cog scale (20-45) where patients are moderately demented, still living at home, but anticipating an accelerated deterioration as part of the natural history of AD, over the next 12 months. Because the primary outcome is efficacy, a triple-blind design is chosen.

25 Several considerations drove our choice of dose. In our previous studies on transgenic mice, doses of 20-30 mg/kg of CQ orally daily for five days per week were markedly effective at inhibiting A β accumulation after 2-3 months of treatment. The human equivalent dose of 1500-2250 mg/day is close to the prescribed antibiotic dose of CQ (600 mg po qid). However, this magnitude of dose, administered for 30 months, would raise concerns about SMON toxicity.

35 As the compound of formula I or II may be conjugated to glucuronide followed by renal excretion, there was some concern that blood levels in the elderly might be elevated by inefficient liver metabolism, constraining the proposed dose even further. Therefore a cautious dose escalation design is chosen, to maximize the chance of detecting a change in outcome measures, while minimizing the risk of adverse effects. The starting dose of 3.3 mg/kg/day, assuming 75 kg average weight, is within

the same order of magnitude of the effective dose in the transgenic mouse model, but only about one tenth of the antibiotic dose.

Since there is no data from the transgenic mouse study of the effectiveness of doses less than 20 mg/kg/day, we reasoned that a beneficial effect might require a longer period of treatment than the 9-12 week duration of the mouse study (Cherny et al., 2001). Therefore a trial length of 36 weeks at an average dose which is approximately one-third of what is effective in the transgenic mice is chosen. The final dose of 10 mg/kg/day is half of an effective dose in mice.

Thirty-six patients will be randomised [18 placebo and 18 (the compound of formula I or II)]. Per protocol analyses will be conducted on 33 patients for analyses up to 24 weeks and 32 patients at 36 weeks. Groups will be similar across most demographic, biological and clinical variables at baseline. The effect of treatment was statistically significant in the more-severely affected group (baseline ADAS-Cog \geq 25), but not the less-severely affected group (ADAS-Cog $<$ 25).

METHODS

Ethical issues: In compliance with Australian Commonwealth and Victorian State laws concerning consent from individuals whose cognitive function may be impaired to the extent of being unable to make informed judgements or decisions, "Consent to Special Procedures" administered by the Victorian Civil and Administrative Tribunal will be obtained for each participant. In addition, third party consent will be obtained from all carers, in accordance with Victoria's Guardianship Laws. SMON will be described in detail in the plain language statement, and discussed verbally with both patient and carer at the time of giving consent. As partially effective treatments are now available for AD, it was considered unethical to have the comparison group on placebo only; hence both treatment groups will be placed on donepezil for the duration of the study. The study was approved by the Royal Melbourne Hospital Research Foundation's Clinical Research and Ethics Committee.

Study population:

Criteria for inclusion in the study were:

- (a) informed consent;
- (b) a diagnosis of probable Alzheimer's disease by NINCDS-ADRDA criteria (McKhann et al., 1984);
- (c) Alzheimer's Disease Assessment Scale-Cognitive (ADAS-Cog) score of 20-45 inclusive (Rosen et al., 1984);

- (d) Mini Mental State Examination (MMSE) score of 10-24 inclusive (Folstein et al., 1975);
- (e) on donepezil hydrochloride 5mg or 10mg for at least 6 months;
- (f) relative or carer willing and able to support the trial;
- 5 (g) able to complete trial examinations; and
- (h) primary sensorial functions intact.

All female patients were postmenopausal.

- Patients are to be excluded if they have a potential allergy to the compound of formula I or II; history or clinical evidence of peripheral neuropathy or
- 10 optic neuropathy; co-existing illnesses or past history which may have affected cognitive function or nerve conduction, including alcohol abuse or dependency; metabolic deficiencies (eg unstable thyroid dysfunction); infections with neurotrophic organisms such as syphilis, HIV, CMV, or EBV; current major depressive episode according to DSM-IV criteria; co-existing illnesses which might confound the adverse
- 15 event profile, such as diabetes, untreated vitamin B12 or folate deficiency, ulcerative colitis, Crohn's disease, chronic diarrhoea, or multiple sclerosis; other co-existing medical conditions which might compromise the patient if s/he were to participate in a clinical trial, such as a neoplasm currently active or likely to recur (except non-melanoma skin cancer), history of immunosuppression, gastrointestinal malabsorption,
- 20 hypertension (BP > 180 mmHg systolic or > 95 mmHg diastolic), cardiac failure (orthopnea, JVP>5cm, or peripheral oedema requiring the prescription of loop diuretics), a history of stroke in the last 6 months or a Hachinski score ≥ 6 , haemoglobin >20% below lower limit of normal range, raised white cell count (20% above reference range), neutropenia (white cell count < 2.5), abnormal liver function tests (>50% above
- 25 reference range), abnormal creatinine clearance (< 75% of reference range), abnormal fasting blood glucose (> 50% above upper limit of normal range), abnormal thyroid function (TSH or T4 >20% outside reference range), or positive hepatitis A,B or C IgM.

- The following factors are to be obtained at baseline to determine whether they correlated with outcome measures: age, sex, premorbid IQ (estimated from Natural
- 30 Adult Reading Text (NART), years of education, serum donepezil hydrochloride, and apolipoprotein E(ApoE) allotype .

- Study design:** The study is a triple blind, placebo-controlled, randomised design. Thirty-six patients and their carers are to be recruited to participate, with patients being
- 35 randomised to receive either the compound of formula I or II or placebo; there will be 18 patients in each arm. The duration of the study was 36 weeks. the compound of

formula I or II dosage was 125mg twice daily from weeks 0-12, increased to 250mg twice daily from weeks 13-24, and finally, 375mg twice daily from weeks 25-36.

5 All patients are to be treated with donepezil hydrochloride for at least 6 months prior to recruitment. The dose of donepezil is optimised by each patient's physician to maximize clinical benefit and to minimize side effects. This dose is maintained for the duration of the study, and patients are to be withdrawn from the study if, at regular review, the dose of donepezil required alteration for any reason.

The study medication and placebo are presented as enteric-coated capsules (125mg were blue, 250mg were brown), randomised in blocks of 6.
10 Presentation after increase to 250mg twice daily was as 2 X 125mg per dose; after increase to 375mg, presentation was twice daily 1 X 125mg and 1 X 250mg per dose. This is to allow the dose to be reduced by 125 mg in each instance, ie to the previous dose, if the patient does not tolerate an increase in dose of study drug or placebo.

15 **Study procedures:** Screening procedures consist of a full medical history, full physical, neurological and ophthalmic examination, blood and urine tests and psychometric tests (ADAS-Cog, MMSE) to confirm the patient's eligibility for the study. Nerve
conduction tests and visual evoked responses are conducted between the screening and baseline visits to provide a baseline measurement, and to exclude patients with
20 undiagnosed peripheral neuropathies or visual disturbances. Blood is collected for ApoE allotyping and determination of baseline plasma levels of the compound of formula I or II, metals and A β prior to randomisation.

The study lasts 36 weeks, with 13 visits (including screening). Eligible subjects we randomised to receive either the compound of formula I or II or placebo.
25 All patients continue their study entry dose of donepezil, and all patients received 100 μ g vitamin B12 IM every four weeks.

Outcome measures: The primary efficacy variable is a change in the baseline score on the Alzheimer's Disease Assessment Scale (ADAS), which is conducted at baseline and
30 at weeks 4, 12, 24 and 36. This readout is chosen for comparability of treatment effects with current therapeutic agents, such as donepezil, for which efficacy trials also used ADAS as their primary outcome measure (Rogers et al., 1998). Although numerous neuropsychological tests could be considered as secondary measures, it is necessary to avoid fatiguing the subjects at review. Therefore the only other cognitive test
35 performed was the Mini-Mental State Exam (MMSE), which is well characterized and easily implemented. The Clinician's Interview Based Impression of Change (CIBIC), a subjective observational index also used in efficacy trials of acetylcholinesterase

inhibitors, is conducted at baseline and at weeks 12, 24 and 36 by an independent researcher who was not part of the study team. Blood samples for measurement of plasma A β and plasma zinc and copper are all taken 4 weekly, by antecubital fossa venepuncture.

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Therapeutic drug monitoring: the compound of formula I or II drug assays are conducted over 6 hours at weeks 12, 24 and 36. The patient's blood is obtained via a heparinized indwelling catheter before the administration of the compound of formula I or II on these days, and then drawn again at 2, 4 and 6 hours post dose. This is done to achieve pharmacokinetic data to correlate with other outcome measures.

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Safety measures: Standard adverse event reporting to a safety monitoring committee, consisting of physicians independent of the study, is conducted to review adverse events at three monthly intervals and on an emergent basis. Following baseline, safety visits are conducted at weeks 2, 4, 8, 12, 16, 20, 24, 26, 28, 32 and 36. The patient and carer are questioned about any changes which might have occurred in the patient's health or medications since the last visit. Standard biochemical, renal and liver function, full blood examination, serum vitamin B12 and folate levels, blood pressure and weight are documented at each visit. A neurological examination is conducted at each visit to assess for peripheral neuropathy and optic neuropathy, and visual evoked responses, nerve conduction studies and a full ophthalmic examination (visual acuity, colour vision, fundal examination and visual field) are conducted at screening, at week 18 and at 2 weeks after trial completion. An ECG was performed at baseline and at weeks 12 and 24.

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Extension study: All patients who completed the Phase II trial are invited to continue on a 48 week, prospective, open-label study of the compound of formula I or II. All are allocated to receive the compound of formula I or II 125mg bid, increased after 2 weeks to 250mg bid, then 375mg BD at 4 weeks, while remaining on donepezil and vitamin B12. Patients who failed to tolerate dose increases beyond the 250mg/day, 500mg/day and 750mg/day dosages in the blinded phase are placed on the highest tolerable dose beyond the previously achieved dose, at the clinical discretion of the investigator in the extension phase. Outcome and safety measures are the same as for the blinded phase. The length of the extension study is based upon an estimation of the time required to complete the blinded Phase II clinical trial, so that subjects will be able to continue to take the drug until they could be advised about the results of the trial.

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Data preparation and statistical analysis: Concealed randomization is conducted in blocks of 6 by the Institute of Drug Technology, a body which is independent of the study. An independent data monitoring company checked for omissions and validated entries in case report forms, and double entered the data into Microsoft Access[®],
5 completing validation and consistency checks. Before the analyses, each patient's randomization arm is labelled either 'A' or 'B'. This ensures that the primary analyses are conducted blind to the subjects' randomization group, and hence were triple blind).

Two-way analysis of variance and covariance is used to analyse the major outcome variables with group (treatment vs placebo) as a between-subjects factor
10 and occasion (baseline vs subsequent measurement occasions) as a within-subjects factor. Evidence for efficacy is indicated by a significant group by occasion interaction. Differences between groups on categorical measures are analysed using exact statistical methods in order to maximise power.

The influence of confounding variables is controlled using analysis of
15 covariance and linear regression models where appropriate. Based on the assumption of 50% shared variance between measurement occasions (i.e. $r=0.70$), power to detect an effect of one standard deviation difference in change between groups from baseline to week 36 would be approximately 80% if 15 subjects were recruited per group. Since an attrition rate of 15% has been observed in similar populations, 18 patients will be
20 recruited into each arm.

The design also includes a subset analysis of outcome measures, in which the cohort is divided into two equal size groups by the median ADAS-Cog score at baseline, yielding a less severely-affected subset, and a more severely-affected subset.

25 It will be apparent to the person skilled in the art that while the invention has been described in some detail for the purposes of clarity and understanding, various modifications and alterations to the embodiments and methods described herein may be made without departing from the scope of the inventive concept disclosed in this specification.

30 References cited in the description and examples are listed on the following pages, and are incorporated herein by this reference.

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25 It will be appreciated by persons skilled in the art that numerous variations and/or modifications may be made to the invention as shown in the specific embodiments without departing from the spirit or scope of the invention as broadly described. The present embodiments are, therefore, to be considered in all respects as illustrative and not restrictive.